

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 9, 2005, 15:07:04 ; Search time 78 Seconds
(without alignments)
639.642 Million cell updates/sec

Title: US-10-001-245C-36

Perfect score: 692

Sequence: 1 DQDVVKDCANHEIKVLVPG.....VLGDNGVLCAIATHAKIRD 129

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A Geneseq_16Dec04:.*
1: Geneseqp1980s:.*
2: Geneseqp1990s:.*
3: Geneseqp2000s:.*
4: Geneseqp2001s:.*
5: Geneseqp2002s:.*
6: Geneseqp2003as:.*
7: Geneseqp2003bs:.*
8: Geneseqp2004s:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	692	100.0	129	5	ABG67011 House dus
2	684	98.8	129	5	ABG67013 House dus
3	684	98.8	129	5	ABG67016 House dus
4	683	98.7	129	5	ABG67015 House dus
5	683	98.7	129	5	ABG67014 House dus
6	682	98.6	129	5	ABG67012 House dus
7	657	94.9	129	5	ABG67019 House dus
8	657	94.9	129	5	ABG67022 House dus
9	656	94.8	129	5	ABG67021 House dus
10	656	94.8	129	5	ABG67020 House dus
11	655	94.7	129	5	ABG67018 House dus
12	655	94.7	129	5	ABG67017 House dus
13	648	93.6	129	5	ABG67010 House dus
14	646	93.4	129	5	ABG66996 House dus
15	643	92.9	129	5	ABG66994 House dus
16	642	92.8	129	5	ABG66993 House dus
17	641	92.6	129	5	ABG66992 House dus
18	641	92.6	129	5	ABG67007 House dus
19	641	92.6	129	5	ABG66976 House dus
20	641	92.6	129	5	ABG67006 House dus
21	641	92.6	129	5	ABG67008 House dus
22	640	92.5	129	5	ABG67001 House dus
23	640	92.5	129	5	ABG67003 House dus
24	639	92.3	129	5	ABG66972 House dus
25	639	92.3	129	5	ABG67000 House dus

26	639	92.3	129	5	ABG66974	House dus
27	639	92.3	129	5	ABG67004	House dus
28	638	92.2	129	5	ABG66995	House dus
29	638	92.2	145	5	ABB76047	Dust mite
30	637	92.1	129	5	ABG67002	House dus
31	637	92.1	129	5	ABG66973	House dus
32	635	91.8	129	5	ABG66997	House dus
33	635	91.8	136	8	ADR87228	Dust mite
34	635	91.8	146	2	AAR39360	Dermatoph
35	635	91.8	146	2	AAR51728	Der p II.
36	635	91.8	146	2	AAW71909	Dermatoph
37	635	91.8	146	2	AAV25581	D. pteron
38	635	91.8	146	2	AAV50357	Dermatoph
39	635	91.8	146	4	AAU18960	House dus
40	635	91.8	146	5	ABG67053	House dus
41	635	91.8	146	6	ABP98483	Amino aci
42	635	91.8	146	7	ADC34831	House dus
43	635	91.8	146	7	ADE38099	European
44	635	91.8	147	2	AAR47064	Protein a
45	633	91.5	129	4	AAU07751	House dus

ALIGNMENTS

RESULT 1
ABG67011
ID ABG67011 standard; protein; 129 AA.
XX
AC ABG67011;
XX
DT 24-SEP-2002 (first entry)
XX
DE House dust mite allergen Der p 2 ALK-G mutant #1.
XX
KW Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
KW vaccine; antiallergic; B cell epitope.
XX
OS Dermatophagoides pteronyssinus.
OS Synthetic.
XX
PN WO200240676-A2.
XX
PD 23-MAY-2002.
XX
PF 16-NOV-2001; 2001WO-DK000764.
XX
PR 16-NOV-2000; 2000DK-00001718.
PR 16-NOV-2000; 2000US-0249361P.
PR 14-JUN-2001; 2001US-0298170P.
XX
PA (ALKA-) ALK-ABELLO AS.
XX
PI Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
XX
DR WPI: 2002-508328/54.
XX
DR N-PSDB; ABK95627.
XX
PT New recombinant mutant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
XX binding affinity.
XX
PS Example 6; Page; 210pp; English.
XX
CC The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the

CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom , and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom ^2 comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IGE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IGE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the
CC specification but was created by the indexer using information in the
CC specification and the corresponding wild-type sequence
XX
SQ Sequence 129 AA;

Query Match 100.0%; Score 692; DB 5; Length 129;
Best Local Similarity 100.0%; Pred. No. 1.3e-73;
Matches 129; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 DQDVKDCANHEIKVELVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQDVKDCANHEIKVELVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
QY 61 LSVDPVGIDPNACHMNCPLVNGQQYDIKTYTNVVKPIAPNSENVVTVKVLGDNGLVACA 120
DB 61 LSVDPVGIDPNACHMNCPLVNGQQYDIKTYTNVVKPIAPNSENVVTVKVLGDNGLVACA 120
QY 121 IATHAKIRD 129
DB 121 IATHAKIRD 129

RESULT 2
ABG67013
ID ABG67013 standard; protein; 129 AA.

XX AC ABG67013;
XX DT 24-SEP-2002 (first entry)
XX DE House dust mite allergen Der p 2 ALK-G mutant #3.
XX DE Immunoglobulin E; IGE; allergen; allergy; mite; hay fever;
KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
KW vaccine; antiallergic; B cell epitope.
OS Dermatophagoides pteronyssinus.
OS Synthetic.
XX WC2002040676-A2.
XX PD 23-MAY-2002.
XX PF 16-NOV-2001; 2001WO-DK000764.
XX PF 16-NOV-2000; 2000DK-00001718.

PR 16-NOV-2000; 2000US-0249361P.
PR 14-JUN-2001; 2001US-0298170P.
XX (ALKA-) ALK-ABELLO AS.
XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
XX WPI; 2002-508328/54.
XX N-PSDB; ABK95629.
XX New recombinant mutant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
PT binding affinity.
XX Example 6; Page; 210pp; English.

CC The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IGE) binding capability of the mutated allergen as compared to the IGE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom , and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom ^2 comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IGE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IGE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the
CC specification but was created by the indexer using information in the
CC specification and the corresponding wild-type sequence
XX
SQ Sequence 129 AA;

Query Match 98.8%; Score 684; DB 5; Length 129;
Best Local Similarity 98.4%; Pred. No. 1.2e-72;
Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 DQDVKDCANHEIKVELVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQDVKDCANHEIKVELVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
QY 61 LSVDPVGIDPNACHMNCPLVNGQQYDIKTYTNVVKPIAPNSENVVTVKVLGDNGLVACA 120
DB 61 LSVDPVGIDPNACHMNCPLVNGQQYDIKTYTNVVKPIAPNSENVVTVKVLGDNGLVACA 120
QY 121 IATHAKIRD 129
DB 121 IATHAKIRD 129

RESULT 3
 ABG67016
 ID ABG67016 standard; protein; 129 AA.
 XX
 AC ABG67016;
 XX
 DT 24-SEP-2002 (first entry)
 XX
 DE House dust mite allergen Der p 2 ALK-G mutant #6.
 XX
 KW Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
 KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
 KW vaccine; antiallergic; B cell epitope.
 XX
 OS Dermatophagoides pteronyssinus.
 XX
 XX Synthetic.
 XX
 PN WO200240676-A2.
 XX
 XX 23-MAY-2002.
 XX
 XX 16-NOV-2001; 2001WO-DK000764.
 XX
 PR 16-NOV-2000; 2000DK-00001718.
 PR 16-NOV-2000; 2000US-0249361P.
 PR 14-JUN-2001; 2001US-0298170P.
 XX
 PA (ALKA-) ALK-ABELLO AS.
 XX
 XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
 XX
 DR WPI; 2002-508328/54.
 DR N-PSDB; ABK95632.
 XX
 PT New recombinant mutant allergen, useful for preventing and/or treating
 PT allergy, comprises multiple mutations and reduced immunoglobulin E
 PT binding affinity.
 XX
 PS Example 6; Page; 210pp; English.
 XX
 CC The invention relates to a recombinant allergen (I) which is a mutant of
 CC a naturally occurring allergen, where the mutant allergen has at least
 CC four primary mutations, which each reduce the specific immunoglobulin E
 CC (IgE) binding capability of the mutated allergen as compared to the IgE
 CC binding capability of the naturally occurring allergen, where each
 CC primary mutation is a substitution of one surface-exposed amino acid
 CC residue with another residue, which does not occur in the same position
 CC in the amino acid sequence of any known homologous protein within the
 CC taxonomic species from which the naturally occurring allergen originates,
 CC and each primary mutation is spaced from each other primary mutation by
 CC at least 15 Angstrom , and the primary mutations are placed in such a
 CC manner that at least one circular surface region with a area of 800
 CC Angstrom² comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IgE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IgE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic

CC anaphylaxis. The present sequence represents a recombinant allergen of
 CC the invention. Note: The present sequence was not shown in the
 CC specification but was created by the indexer using information in the
 CC specification and the corresponding wild-type sequence
 XX
 SQ Sequence 129 AA;
 Query Match 98.8%; Score 684; DB 5; Length 129;
 Best Local Similarity 98.4%; Pred. No. 1.2e-72;
 Matches 127; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 QY 1 DQVDVKDCANHEIKVLPFGCHGNPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
 DB 1 DQVDVKDCANHEIKVLPFGCHGNPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
 QY 61 LSVDFVGIDPNACHYNNCPVNGQQYDIKYTNWPKIAPNSENVVTVKVLGNGVLACA 120
 DB 61 LSVDFVGIDPNACHYNNCPVNGQQYDIKYTNWPKIAPNSENVVTVKVLGNGVLACA 120
 QY 121 IATHAKIRD 129
 DB 121 IATHAKIQD 129
 RESULT 4
 ABG67015
 ID ABG67015 standard; protein; 129 AA.
 XX
 AC ABG67015;
 XX
 DT 24-SEP-2002 (first entry)
 XX
 DE House dust mite allergen Der p 2 ALK-G mutant #5.
 XX
 KW Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
 KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
 KW vaccine; antiallergic; B cell epitope.
 XX
 OS Dermatophagoides pteronyssinus.
 XX
 XX Synthetic.
 XX
 PN WO200240676-A2.
 XX
 XX 23-MAY-2002.
 XX
 XX 16-NOV-2001; 2001WO-DK000764.
 XX
 PR 16-NOV-2000; 2000DK-00001718.
 PR 16-NOV-2000; 2000US-0249361P.
 PR 14-JUN-2001; 2001US-0298170P.
 XX
 PA (ALKA-) ALK-ABELLO AS.
 XX
 XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
 XX
 DR WPI; 2002-508328/54.
 DR N-PSDB; ABK95631.
 XX
 PT New recombinant mutant allergen, useful for preventing and/or treating
 PT allergy, comprises multiple mutations and reduced immunoglobulin E
 PT binding affinity.
 XX
 PS Example 6; Page; 210pp; English.
 XX
 CC The invention relates to a recombinant allergen (I) which is a mutant of
 CC a naturally occurring allergen, where the mutant allergen has at least
 CC four primary mutations, which each reduce the specific immunoglobulin E
 CC (IgE) binding capability of the mutated allergen as compared to the IgE
 CC binding capability of the naturally occurring allergen, where each
 CC primary mutation is a substitution of one surface-exposed amino acid
 CC residue with another residue, which does not occur in the same position
 CC in the amino acid sequence of any known homologous protein within the
 CC taxonomic species from which the naturally occurring allergen originates,
 CC and each primary mutation is spaced from each other primary mutation by
 CC at least 15 Angstrom , and the primary mutations are placed in such a
 CC manner that at least one circular surface region with a area of 800
 CC Angstrom² comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IgE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IgE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic

CC and each primary mutation is spaced from each other primary mutation by
 CC at least 15 Angstrom , and the primary mutations are placed in such a
 CC manner that at least one circular surface region with a area of 800
 CC Angstrom² comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IGE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IGE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
 CC anaphylaxis. The present sequence represents a recombinant allergen of
 CC the invention. Note: The present sequence was not shown in the
 CC specification but was created by the indexer using information in the
 CC specification and the corresponding wild-type sequence

XX Sequence 129 AA;

Query Match 98.7%; Score 683; DB 5; Length 129;
 Best Local Similarity 98.4%; Pred. No. 1.6e-72;
 Matches 127; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
 DB 1 DQVDVKDCANHEIKVLPVCGHSEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

QY 61 LSVDPFGIDPNACHYMNCPVLVNGQQYDIKYTNWVPKIPNSENVVTVKVLGDNGLACA 120
 DB 61 LSVDPFGIDPNACHYMNCPVLVNGQQYDIKYTNWVPKIPNSENVVTVKVLGDNGLACA 120

QY 121 IATHAKIRD 129
 DB 121 IATHAKIQD 129

RESULT 5
 ABG67014
 ID ABG67014 standard; protein; 129 AA.

XX ABG67014;

XX 24-SEP-2002 (first entry)

DE House dust mite allergen Der p 2 ALK-G mutant #4.

XX Immunoglobulin E; IGE; allergen; allergy; mite; hay fever;
 KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
 KW vaccine; antiallergic; B cell epitope.

XX Dermatoaphagoides pteronyssinus.
 OS Synthetic.

XX WO200240676-A2.

XX 23-MAY-2002.

XX 16-NOV-2001; 2001WO-DK000764.

XX 16-NOV-2000; 2000DK-00001718.

PR 16-NOV-2000; 2000US-0249361P.

PR 14-JUN-2001; 2001US-0298170P.

XX (ALKA-) ALK-ABELLO AS.

XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;

XX WPI; 2002-508328/54.

DR N-PSDB; ABK95630.

XX New recombinant mutant allergen, useful for preventing and/or treating
 PT allergy, comprises multiple mutations and reduced immunoglobulin E
 PT binding affinity.

XX Example 6; Page; 210pp; English.

XX The invention relates to a recombinant allergen (I) which is a mutant of
 CC a naturally occurring allergen, where the mutant allergen has at least
 CC four primary mutations, which each reduce the specific immunoglobulin E
 CC (IGE) binding capability of the mutated allergen as compared to the IGE
 CC binding capability of the naturally occurring allergen, where each
 CC primary mutation is a substitution of one surface-exposed amino acid
 CC residue with another residue, which does not occur in the same position
 CC in the amino acid sequence of any known homologous protein within the
 CC taxonomic species from which the naturally occurring allergen originates,
 CC and each primary mutation is spaced from each other primary mutation by
 CC at least 15 Angstrom , and the primary mutations are placed in such a
 CC manner that at least one circular surface region with a area of 800
 CC Angstrom² comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IGE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IGE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
 CC anaphylaxis. The present sequence represents a recombinant allergen of
 CC the invention. Note: The present sequence was not shown in the
 CC specification but was created by the indexer using information in the
 CC specification and the corresponding wild-type sequence

XX Sequence 129 AA;

Query Match 98.7%; Score 683; DB 5; Length 129;

Best Local Similarity 98.4%; Pred. No. 1.6e-72;

Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

DB 1 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

QY 61 LSVDPFGIDPNACHYMNCPVLVNGQQYDIKYTNWVPKIPNSENVVTVKVLGDNGLACA 120

DB 61 LSVDPFGIDPNACHYMNCPVLVNGQQYDIKYTNWVPKIPNSENVVTVKVLGDNGLACA 120

QY 121 IATHAKIRD 129

DB 121 IATHAKIQD 129

RESULT 6

ABG67012
ID ABG67012 standard; protein; 129 AA.
XX AC ABG67012;
XX DT 24-SEP-2002 (first entry)
XX DE House dust mite allergen Der p 2 ALK-G mutant #2.
XX KW Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
XX KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
XX KW vaccine; antiallergic; B cell epitope.
XX OS Dermatophagoides pteronyssinus.
XX OS Synthetic.
XX PN W0200240676-A2.
XX PD 23-MAY-2002.
XX PF 16-NOV-2001; 2001WO-DK000764.
XX PR 16-NOV-2000; 2000DK-00001718.
XX PR 16-NOV-2000; 2000US-0249361P.
XX PR 14-JUN-2001; 2001US-0298170P.
XX PA (ALKA-) ALK-ABELLO AS.
XX PI Holm J, Ipeen H, Nedergaard Larsen J, Spangfort MD;
XX WPI; 2002-508328/54.
XX DR N-PSDB; ABK95628.
XX PT New recombinant mutant allergen, useful for preventing and/or treating
XX PT allergy, comprises multiple mutations and reduced immunoglobulin E
XX PT binding affinity.
XX PS Example 6; Page; 210pp; English.
XX CC The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom, and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom² comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridizes to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IgE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IgE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of

CC the invention. Note: The present sequence was not shown in the
CC specification but was created by the indexer using information in the
CC specification and the corresponding wild-type sequence
XX SQ Sequence 129 AA;
Query Match 98.6%; Score 682; DB 5; Length 129;
Best Local Similarity 98.4%; Pred. No. 2.1e-72;
Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 DQVDVKDCANHEIKELVFGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQVDVKDCANHEIKELVFGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
QY 61 LSVDFVPGIDPNACHYMNCPVNGQQYDIKYTNVVKPIAPNSNVVTVKVLGNGVLACA 120
DB 61 LSVDFVPGIDPNACHYMNCPVNGQQYDIKYTNVVKPIAPNSNVVTVKVLGNGVLACA 120
QY 121 IATHAKIRD 129
DB 121 IATHAKIQD 129
RESULT 7
ABG67019
ID ABG67019 standard; protein; 129 AA.
XX AC ABG67019;
XX DT 24-SEP-2002 (first entry)
XX DE House dust mite allergen Der p 2 ALK-G mutant #9.
XX KW Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
XX KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
XX KW vaccine; antiallergic; B cell epitope.
XX OS Dermatophagoides pteronyssinus.
XX OS Synthetic.
XX PN W0200240676-A2.
XX PD 23-MAY-2002.
XX PF 16-NOV-2001; 2001WO-DK000764.
XX PR 16-NOV-2000; 2000DK-00001718.
XX PR 16-NOV-2000; 2000US-0249361P.
XX PR 14-JUN-2001; 2001US-0298170P.
XX PA (ALKA-) ALK-ABELLO AS.
XX PI Holm J, Ipeen H, Nedergaard Larsen J, Spangfort MD;
XX WPI; 2002-508328/54.
XX DR N-PSDB; ABK95635.
XX PT New recombinant mutant allergen, useful for preventing and/or treating
XX PT allergy, comprises multiple mutations and reduced immunoglobulin E
XX PT binding affinity.
XX PS Example 6; Page; 210pp; English.
XX CC The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom, and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom² comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridizes to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IgE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IgE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of

at least 15 Angstrom , and the primary mutations are placed in such a manner that at least one circular surface region with a area of 800 Angstrom ^2 comprises no mutation. Also included are a composition comprising two or more of the recombinant allergens, where the variant allergen is defined by having at least one primary mutation, which is absent in at least one of the other variants, and for each variant no secondary mutation is present within a radius of 15 Angstrom from each absent primary mutation; a DNA sequence encoding the recombinant allergen or its derivative, partial sequence or degenerated sequence, or a sequence which hybridises to it under stringent conditions, where the derivative, partial sequence, degenerated sequence or hybridising sequence encodes a peptide having at least one B cell epitope; an expression vector comprising the DNA and a host cell comprising the vector. The recombinant allergen is useful as a pharmaceutical, for preparing a pharmaceutical for preventing and/or treating allergy, or in a diagnostic assay for assessing relevance, safety or outcome of therapy of a subject, where an IGE containing sample of the subject is mixed with the recombinant allergen and assessed for the level of reactivity between the IGE in the sample and the recombinant allergen. The recombinant allergen or compositions are useful for generating an immune response in a subject, for vaccination or treatment of a subject or for the treatment, prevention or alleviation of allergic reactions in a subject e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic anaphylaxis. The present sequence represents a recombinant allergen of the invention. Note: The present sequence was not shown in the specification but was created by the indexer using information in the specification and the corresponding wild-type sequence

SQ Sequence 129 AA;

Query Match 94.9%; Score 657; DB 5; Length 129;
Best Local Similarity 95.3%; Pred. No. 1.9e-69;
Matches 123; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
QY 61 LSVDPVPGIDPNACHYMNCPVLNGQQYDIKYTNVNPKIAPNSENVVTVKVLGDNGLACA 120
DB 61 LSVDPVPGIDPNACHYMNCPVLNGQQYDIKYTNVNPKIAPNSENVVTVKVLGDNGLACA 120
QY 121 IATHAKIRD 129
DB 121 IATHAKIQD 129

RESULT 8

ABG67022

ID ABG67022 standard; protein; 129 AA.

AC ABG67022;

XX 24-SEP-2002 (first entry)

XX House dust mite allergen Der p 2 ALK-G mutant #12.

XX Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;

KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;

XX vaccine; anti-allergic; B cell epitope.

OS Dermatophagoides pteronyssinus.

OS Synthetic.

XX WO200240676-A2.

PN 23-MAY-2002.

XX 16-NOV-2001; 2001WO-DK000764.

XX 16-NOV-2000; 2000DK-00001718.

PR 16-NOV-2000; 2000US-0249361P.

PR 14-JUN-2001; 2001US-0298170P.

XX (ALKA-) ALK-ABELLO AS.
PA Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
XX WPI; 2002-508328/54.
DR N-PSDB; ABK95638.
XX New recombinant mutant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
PT binding affinity.
XX Example 6; Page; 210pp; English.

XX The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IGE) binding capability of the mutated allergen as compared to the IGE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom , and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom ^2 comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IGE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IGE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the
CC specification but was created by the indexer using information in the
CC specification and the corresponding wild-type sequence

XX Sequence 129 AA;

Query Match 94.9%; Score 657; DB 5; Length 129;

Best Local Similarity 95.3%; Pred. No. 1.9e-69;

Matches 123; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

DB 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

QY 61 LSVDPVPGIDPNACHYMNCPVLNGQQYDIKYTNVNPKIAPNSENVVTVKVLGDNGLACA 120

DB 61 LSVDPVPGIDPNACHYMNCPVLNGQQYDIKYTNVNPKIAPNSENVVTVKVLGDNGLACA 120

QY 121 IATHAKIRD 129

DB 121 IATHAKIQD 129

RESULT 9

ABG67021

ID XX ABG67021 standard; protein; 129 AA.
AC XX ABG67021;
DT XX 24-SEP-2002 (first entry)
DE XX House dust mite allergen Der p 2 ALK-G mutant #11.
XX XX Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
KW vaccine; antiallergic; B cell epitope.
XX XX Dermatophagoides pteronyssinus.
OS Synthetic.
XX WO200240676-A2.
XX 23-MAY-2002.
XX 16-NOV-2001; 2001WO-DK000764.
XX 16-NOV-2000; 2000DK-00001718.
XX 16-NOV-2000; 2000US-0249361P.
XX 14-JUN-2001; 2001US-0298170P.
XX (ALKA-) ALK-ABELLO AS.
XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
PI WPI; 2002-508328/54.
DR N-PSDB; ABK95637.
XX New recombinant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
PT binding affinity.
XX Example 6; Page; 210pp; English.
XX The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom, and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom² comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IgE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IgE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the

CC specification but was created by the indexer using information in the
CC specification and the corresponding wild-type sequence
XX Sequence 129 AA;
SQ Query Match 94.8%; Score 656; DB 5; Length 129;
Best Local Similarity 95.3%; Pred. No. 2.5e-69;
Matches 123; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
QY 1 DQVDVKDCANHEIKVELVPGCHGNEPCITGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQVDVKDCANHEIKVELVPGCHGSEPCITHSRGKPFQLEALFEANQNSATAKIEIKASIDG 60
QY 61 LSVDFPGIDPNACHYMNCPVNGQQYDIKTYMNPVKIAPNSNVVTVKVLGNGVLACA 120
DB 61 LSVDFPGIDPNACHYMNCPVNGQQYDIKTYMNPVKIAPNSNVVTVKVLGNGVLACA 120
QY 121 IATHAKIRD 129
DB 121 IATHAKIQD 129
RESULT 10
ABG67020
ID ABG67020 standard; protein; 129 AA.
XX AC ABG67020;
XX 24-SEP-2002 (first entry)
DT House dust mite allergen Der p 2 ALK-G mutant #10.
DE rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
KW vaccine; antiallergic; B cell epitope.
XX Dermatophagoides pteronyssinus.
OS Synthetic.
XX WO200240676-A2.
XX 23-MAY-2002.
XX 16-NOV-2001; 2001WO-DK000764.
XX 16-NOV-2000; 2000DK-00001718.
XX 16-NOV-2000; 2000US-0249361P.
XX 14-JUN-2001; 2001US-0298170P.
XX (ALKA-) ALK-ABELLO AS.
XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
PI WPI; 2002-508328/54.
DR N-PSDB; ABK95636.
XX New recombinant mutant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
PT binding affinity.
XX Example 6; Page; 210pp; English.
XX The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom, and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom² comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IgE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IgE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the

CC manner that at least one circular surface region with a area of 800
 CC Angstrom ^2 comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IGE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IGE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
 CC anaphylaxis. The present sequence represents a recombinant allergen of
 CC the invention. Note: The present sequence was not shown in the
 CC specification but was created by the indexer using information in the
 CC specification and the corresponding wild-type sequence

XX Sequence 129 AA;

Query Match 94.8%; Score 656; DB 5; Length 129;
 Best Local Similarity 95.3%; Pred. No. 2.5e-69;
 Matches 123; Conservative 2; Mismatches 4; Indels 0; Gaps 0;
 QY 1 DQVDVKDCANHEIKVELVPGCHGNEPCIIIGRKPFOLEALFEANQNSATAKIEIKASIDG 60
 DB 1 DQVDVKDCANHEIKVELVPGCHGNEPCIIHSGRKPFQLEALFEANQNSATAKIEIKASIDG 60
 QY 61 LSVDPVPGIDPNACHYMCPLVNGQQYDIKYTNWVVKIAPNSENVVTVKVLGDNGLACA 120
 DB 61 LSVDPVPGIDPNACHYMCPLVNGQQYDIKYTNWVVKIAPNSENVVTVKVLGDNGLACA 120
 QY 121 IATHAKIRD 129
 DB 121 IATHAKIQD 129

RESULT 11

ABG67018

ID ABG67018 standard; protein; 129 AA.

AC ABG67018;

XX 24-SEP-2002 (first entry)

XX House dust mite allergen Der p 2 ALK-G mutant #8.

XX Immunoglobulin E; IgE; allergen; allergy; mite; hay fever;
 KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
 KW vaccine; anti-allergic; B cell epitope.

XX Dermatophagoides pteronyssinus.

OS Synthetic.

XX WO200240676-A2.

XX 23-MAY-2002.

XX 16-NOV-2001; 2001WO-DK000764.

XX 16-NOV-2000; 2000DK-00001718.

PR 16-NOV-2000; 2000US-0249361P.

PR 14-JUN-2001; 2001US-0298170P.

XX

PA (ALKA-) ALK-ABELLO AS.
 XX Holm J, Ipeen H, Nedergaard Larsen J, Spangfort MD;
 PI WPI; 2002-508328/54.
 XX N-PSDB; ABK95634.
 DR New recombinant mutant allergen, useful for preventing and/or treating
 XX allergy, comprises multiple mutations and reduced immunoglobulin E
 PT binding affinity.
 PT Example 6; Page; 210pp; English.
 XX The invention relates to a recombinant allergen (I) which is a mutant of
 CC a naturally occurring allergen, where the mutant allergen has at least
 CC four primary mutations, which each reduce the specific immunoglobulin E
 CC (IGE) binding capability of the mutated allergen as compared to the IGE
 CC binding capability of the naturally occurring allergen, where each
 CC primary mutation is a substitution of one surface-exposed amino acid
 CC residue with another residue, which does not occur in the same position
 CC in the amino acid sequence of any known homologous protein within the
 CC taxonomic species from which the naturally occurring allergen originates,
 CC and each primary mutation is spaced from each other primary mutation by
 CC at least 15 Angstrom , and the primary mutations are placed in such a
 CC manner that at least one circular surface region with a area of 800
 CC Angstrom ^2 comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IGE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IGE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
 CC anaphylaxis. The present sequence represents a recombinant allergen of
 CC the invention. Note: The present sequence was not shown in the
 CC specification but was created by the indexer using information in the
 CC specification and the corresponding wild-type sequence

XX Sequence 129 AA;

Query Match 94.7%; Score 655; DB 5; Length 129;
 Best Local Similarity 95.3%; Pred. No. 3.3e-69;
 Matches 123; Conservative 2; Mismatches 4; Indels 0; Gaps 0;
 QY 1 DQVDVKDCANHEIKVELVPGCHGNEPCIIIGRKPFOLEALFEANQNSATAKIEIKASIDG 60
 DB 1 DQVDVKDCANHEIKVELVPGCHGNEPCIIHSGRKPFQLEALFEANQNSATAKIEIKASIDG 60
 QY 61 LSVDPVPGIDPNACHYMCPLVNGQQYDIKYTNWVVKIAPNSENVVTVKVLGDNGLACA 120
 DB 61 LSVDPVPGIDPNACHYMCPLVNGQQYDIKYTNWVVKIAPNSENVVTVKVLGDNGLACA 120
 QY 121 IATHAKIRD 129
 DB 121 IATHAKIQD 129

RESULT 12

ABG67017

ID ABG67017 standard; protein; 129 AA.

XX ABG67017;
XX 24-SEP-2002 (first entry)
XX House dust mite allergen Der p 2 ALK-G mutant #7.
XX Immunoglobulin E; IgE; allergen; allergy; mutein; hay fever;
XX rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; mutant;
XX vaccine; antiallergic; B cell epitope.
XX
XX Dermatophagoides pteronyssinus.
OS Synthetic.
XX
XX WO200240676-A2.
XX
XX 23-MAY-2002.
XX
XX 16-NOV-2001; 2001WO-DK000764.
XX
XX 16-NOV-2000; 2000DK-00001718.
XX 16-NOV-2000; 2000US-0249361P.
XX 14-JUN-2001; 2001US-0298170P.
XX
XX (ALKA-) ALK-ABELLO AS.
XX
XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
XX
XX WPI; 2002-508328/54.
XX N-PSDB; ABK95633.
XX
XX New recombinant mutant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
PT binding affinity.
XX
XX Example 6; Page; 210pp; English.
XX
XX The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom , and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom ^2 comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IgE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IgE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the
CC specification but was created by the indexer using information in the

CC specification and the corresponding wild-type sequence
XX
XX Sequence 129 AA;
XX
XX Query Match 94.7%; Score 655; DB 5; Length 129;
XX Best Local Similarity 95.3%; Pred. NO. 3.3e-69;
XX Matches 123; Conservative 2; Mismatches 4; Indels 0; Gaps 0;
QY 1 DQVDVDCANHEIKEVLPFGCHGNBPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQVDVDCANHEIKEVLPFGCHGNBPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
QY 61 LSVDPFGIDPNACHNCPVNGQQYDIKYTNVNPKIAPNSNVVTVKVLGNGVLACA 120
DB 61 LSVDPFGIDPNACHNCPVNGQQYDIKYTNVNPKIAPNSNVVTVKVLGNGVLACA 120
QY 121 IATHAKIRD 129
DB 121 IATHAKIQD 129
RESULT 13
ABG67010
ID ABG67010 standard; protein; 129 AA.
XX
XX ABG67010;
XX
XX 24-SEP-2002 (first entry)
XX House dust mite allergen Der p 2 ALK-G.
XX
XX Immunoglobulin E; IgE; allergen; allergy; hay fever; rhinoconjunctivitis;
XX rhinitis; asthma; systemic anaphylaxis; vaccine; antiallergic;
XX B cell epitope.
XX
XX Dermatophagoides pteronyssinus.
XX
XX WO200240676-A2.
XX
XX 23-MAY-2002.
XX
XX 16-NOV-2001; 2001WO-DK000764.
XX
XX 16-NOV-2000; 2000DK-00001718.
XX 16-NOV-2000; 2000US-0249361P.
XX 14-JUN-2001; 2001US-0298170P.
XX
XX (ALKA-) ALK-ABELLO AS.
XX
XX Holm J, Ipsen H, Nedergaard Larsen J, Spangfort MD;
XX
XX WPI; 2002-508328/54.
XX N-PSDB; ABK95626.
XX
XX New recombinant mutant allergen, useful for preventing and/or treating
PT allergy, comprises multiple mutations and reduced immunoglobulin E
PT binding affinity.
XX
XX Example 6; Page 96; 210pp; English.
XX
XX The invention relates to a recombinant allergen (I) which is a mutant of
CC a naturally occurring allergen, where the mutant allergen has at least
CC four primary mutations, which each reduce the specific immunoglobulin E
CC (IgE) binding capability of the mutated allergen as compared to the IgE
CC binding capability of the naturally occurring allergen, where each
CC primary mutation is a substitution of one surface-exposed amino acid
CC residue with another residue, which does not occur in the same position
CC in the amino acid sequence of any known homologous protein within the
CC taxonomic species from which the naturally occurring allergen originates,
CC and each primary mutation is spaced from each other primary mutation by
CC at least 15 Angstrom , and the primary mutations are placed in such a
CC manner that at least one circular surface region with a area of 800
CC Angstrom ^2 comprises no mutation. Also included are a composition
CC comprising two or more of the recombinant allergens, where the variant
CC allergen is defined by having at least one primary mutation, which is
CC absent in at least one of the other variants, and for each variant no
CC secondary mutation is present within a radius of 15 Angstrom from each
CC absent primary mutation; a DNA sequence encoding the recombinant allergen
CC or its derivative, partial sequence or degenerated sequence, or a
CC sequence which hybridises to it under stringent conditions, where the
CC derivative, partial sequence, degenerated sequence or hybridising
CC sequence encodes a peptide having at least one B cell epitope; an
CC expression vector comprising the DNA and a host cell comprising the
CC vector. The recombinant allergen is useful as a pharmaceutical, for
CC preparing a pharmaceutical for preventing and/or treating allergy, or in
CC a diagnostic assay for assessing relevance, safety or outcome of therapy
CC of a subject, where an IgE containing sample of the subject is mixed with
CC the recombinant allergen and assessed for the level of reactivity between
CC the IgE in the sample and the recombinant allergen. The recombinant
CC allergen or compositions are useful for generating an immune response in
CC a subject, for vaccination or treatment of a subject or for the
CC treatment, prevention or alleviation of allergic reactions in a subject
CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
CC anaphylaxis. The present sequence represents a recombinant allergen of
CC the invention. Note: The present sequence was not shown in the
CC specification but was created by the indexer using information in the

CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IGE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IGE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
 CC anaphylaxis. The present sequence represents a wild-type allergen of the
 CC invention
 XX
 SQ Sequence 129 AA;

Query Match 93.6%; Score 648; DB 5; Length 129;
 Best Local Similarity 93.8%; Pred. No. 2.2e-68;
 Matches 121; Conservative 2; Mismatches 6; Indels 0; Gaps 0;
 QY 1 DQDVVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
 Db 1 DQDVVKDCANHEIKKVLVPGCHGSEPCIIHRGKPFQLEALFEANQNSKATAKIEIKASIDG 60
 QY 61 LSVDPFGIDPNACHYMNCPVLNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGVLA 120
 Db 61 LEVDVPGIDPNACHYMKCPLVKGQQYDIKYTNVVPKIAPKSENVVTVKVLGDNGVLA 120
 QY 121 IATHAKIRD 129
 Db 121 IATHAKIRD 129

RESULT 14

ABG66996
 ID ABG66996 standard; protein; 129 AA.

XX
 AC ABG66996;

XX
 DT 24-SRP-2002 (first entry)

XX
 DE House dust mite allergen Der p 2 isoform ALK-120.

XX
 KW Immunoglobulin E; IgE; allergen; allergy; hay fever; house dust mite;
 KW rhinoconjunctivitis; rhinitis; asthma; systemic anaphylaxis; isoform;
 KW vaccine; anti-allergic; B cell epitope; Der p 2.

XX
 OS Dermatophagoides pteronyssinus.

XX
 PN WO200240676-A2.

XX
 PD 23-MAY-2002.

XX
 PF 16-NOV-2001; 2001WO-DK000764.

XX
 PR 16-NOV-2000; 2000DK-00001718.

XX
 PR 16-NOV-2000; 2000US-0249361P.

XX
 PR 14-JUN-2001; 2001US-0298170P.

XX
 PA (ALKA-) ALK-ABELLO AS.

XX
 PI Holm J, Ipeen H, Nedergaard Larsen J, Spangfort MD;

XX
 XX WPI; 2002-508328/54.

XX
 PT New recombinant mutant allergen, useful for preventing and/or treating
 PT allergy, comprises multiple mutations and reduced immunoglobulin E
 PT binding affinity.

XX
 PS Example 5; Page; 210pp; English.

XX
 CC The invention relates to a recombinant allergen (I) which is a mutant of
 CC a naturally occurring allergen, where the mutant allergen has at least
 CC four primary mutations, which each reduce the specific immunoglobulin E
 CC (IGE) binding capability of the mutated allergen as compared to the IGE
 CC binding capability of the naturally occurring allergen, where each
 CC primary mutation is a substitution of one surface-exposed amino acid
 CC residue with another residue, which does not occur in the same position
 CC in the amino acid sequence of any known homologous protein within the
 CC taxonomic species from which the naturally occurring allergen originates,
 CC and each primary mutation is spaced from each other primary mutation by
 CC at least 15 Angstrom , and the primary mutations are placed in such a
 CC manner that at least one circular surface region with a area of 800
 CC Angstrom ^2 comprises no mutation. Also included are a composition
 CC comprising two or more of the recombinant allergens, where the variant
 CC allergen is defined by having at least one primary mutation, which is
 CC absent in at least one of the other variants, and for each variant no
 CC secondary mutation is present within a radius of 15 Angstrom from each
 CC absent primary mutation; a DNA sequence encoding the recombinant allergen
 CC or its derivative, partial sequence or degenerated sequence, or a
 CC sequence which hybridises to it under stringent conditions, where the
 CC derivative, partial sequence, degenerated sequence or hybridising
 CC sequence encodes a peptide having at least one B cell epitope; an
 CC expression vector comprising the DNA and a host cell comprising the
 CC vector. The recombinant allergen is useful as a pharmaceutical, for
 CC preparing a pharmaceutical for preventing and/or treating allergy, or in
 CC a diagnostic assay for assessing relevance, safety or outcome of therapy
 CC of a subject, where an IGE containing sample of the subject is mixed with
 CC the recombinant allergen and assessed for the level of reactivity between
 CC the IGE in the sample and the recombinant allergen. The recombinant
 CC allergen or compositions are useful for generating an immune response in
 CC a subject, for vaccination or treatment of a subject or for the
 CC treatment, prevention or alleviation of allergic reactions in a subject
 CC e.g. hay fever, rhinoconjunctivitis, rhinitis, asthma or systemic
 CC anaphylaxis. The present sequence represents an isoform of the house dust
 CC mite allergen Der p 2
 XX
 SQ Sequence 129 AA;

Query Match 93.4%; Score 646; DB 5; Length 129;
 Best Local Similarity 93.0%; Pred. No. 3.9e-68;
 Matches 120; Conservative 3; Mismatches 6; Indels 0; Gaps 0;

QY 1 DQDVVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
 Db 1 DQDVVKDCANHEIKKVLVPGCHGSEPCIIHRGKPFQLEALFEANQNSKATAKIEIKASIDG 60
 QY 61 LSVDPFGIDPNACHYMNCPVLNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGVLA 120
 Db 61 LEVDVPGIDPNACHYMKCPLVKGQQYDIKYTNVVPKIAPKSENVVTVKVLGDNGVLA 120
 QY 121 IATHAKIRD 129
 Db 121 IATHAKIRD 129

RESULT 15

ABG66994

ID ABG66994 standard; protein; 129 AA.

XX
 AC ABG66994;

XX
 DT 24-SEP-2002 (first entry)

XX
 DE House dust mite allergen Der p 2 isoform ALK-104.

XX
 KW Immunoglobulin E; IgE; allergen; allergy; hay fever; house dust mite;

KW rhinoconductivitis; rhinitis; asthma; systemic anaphylaxis; isoform;
XX vaccine; antiallergic; B cell epitope; Der p 2.
OS Dermatophagoides pteronyssinus.
XX WO200240676-A2.
XX 23-MAY-2002.
XX
XX 16-NOV-2001; 2001WO-DK000764.
XX 16-NOV-2000; 2000DK-00001718.
XX 16-NOV-2000; 2000US-0249361P.
XX 14-JUN-2001; 2001US-0298170P.
XX
XX (ALKA-) ALK-ABELLO AS.
XX
XX Holm J, Ipeen H, Nedergaard Larsen J, Spangfort MD;
XX WPI; 2002-508328/54.
XX
XX New recombinant mutant allergen, useful for preventing and/or treating
XX allergy, comprises multiple mutations and reduced immunoglobulin E
XX binding affinity.
XX
XX Example 5; Page; 210pp; English.
XX
XX The invention relates to a recombinant allergen (I) which is a mutant of
XX a naturally occurring allergen, where the mutant allergen has at least
XX four primary mutations, which each reduce the specific immunoglobulin E
XX (IgE) binding capability of the mutated allergen as compared to the IgE
XX binding capability of the naturally occurring allergen, where each
XX primary mutation is a substitution of one surface-exposed amino acid
XX residue with another residue, which does not occur in the same position
XX in the amino acid sequence of any known homologous protein within the
XX taxonomic species from which the naturally occurring allergen originates,
XX and each primary mutation is spaced from each other primary mutation by
XX at least 15 Angstrom , and the primary mutations are placed in such a
XX manner that at least one circular surface region with a area of 800
XX Angstrom ^2 comprises no mutation. Also included are a composition
XX comprising two or more of the recombinant allergens, where the variant
XX allergen is defined by having at least one primary mutation, which is
XX absent in at least one of the other variants, and for each variant no
XX secondary mutation is present within a radius of 15 Angstrom from each
XX absent primary mutation; a DNA sequence encoding the recombinant allergen
XX or its derivative, partial sequence or degenerated sequence, or a
XX sequence which hybridises to it under stringent conditions, where the
XX derivative, partial sequence, degenerated sequence or hybridising
XX sequence encodes a peptide having at least one B cell epitope; an
XX expression vector comprising the DNA and a host cell comprising the
XX vector. The recombinant allergen is useful as a pharmaceutical, for
XX preparing a pharmaceutical for preventing and/or treating allergy, or in
XX a diagnostic assay for assessing relevance, safety or outcome of therapy
XX of a subject, where an IgE containing sample of the subject is mixed with
XX the recombinant allergen and assessed for the level of reactivity between
XX the IgE in the sample and the recombinant allergen. The recombinant
XX allergen or compositions are useful for generating an immune response in
XX a subject, for vaccination or treatment of a subject or for the
XX treatment, prevention or alleviation of allergic reactions in a subject
XX e.g. hay fever, rhinoconductivitis, rhinitis, asthma or systemic
XX anaphylaxis. The present sequence represents an isoform of the house dust
XX mite allergen Der p 2
XX
XX Sequence 129 AA;

Query Match 92.9%; Score 643; DB 5; Length 129;
Best Local Similarity 92.2%; Pred. No. 8.e-68;
Matches 119; Conservative 4; Mismatches 6; Indels 0; Gaps 0;
QY 1 DQVDVKDCANHEIKVLVPCGKGPFCLEALFEANQNSATAKIEIKASIDG 60
DB 1 DQVDVKDCANHEIKVLVPCGKGPFCLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDFGIDPNACHYMNCPVLVNGQQYDIKTYTNVPKIAPNSNVVVTKVLGNGVLACA 120
Db 61 LEVDVPGIDPNACHYMKCPVLVKGQQYDIKTYTNVPKIAPKSNVVVTKVIGDNGVLACA 120
Qy 121 IATHAKIRD 129
Db 121 IATHAKIRD 129
Search completed: September 9, 2005, 15:21:35
Job time : 80 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 9, 2005, 15:10:19 ; Search time 24 Seconds
(without alignments)
517.165 Million cell updates/sec

Title: US-10-001-245C-36

Perfect score: 692

Sequence: 1 DQVDVKDCANHEIKVELVPG.....VLGDNGVLACAIATHAKIRD 129

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 79: *
1: pir1: *
2: pir2: *
3: pir3: *
4: pir4: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	635	91.8	146	2 A60381	allergen Der p II
2	592	85.5	129	2 JU0394	allergen Der f II
3	590	85.3	138	2 B61241	allergen Der f II
4	588	85.0	138	2 A61241	allergen Der f II
5	585	84.5	129	2 A61501	allergen Der f II
6	240.5	34.8	141	2 S66500	allergen Lep d 1.0
7	228.5	33.0	141	2 S66499	allergen Lep d 1.0
8	112	16.2	151	2 I53929	epididymal secreto
9	112	16.2	151	2 I38365	epididymal secreto
10	104.5	15.1	149	2 I69229	epididymal secreto
11	95.5	13.8	186	2 T32408	hypothetical prote
12	84.5	12.2	408	2 G83893	hypothetical prote
13	83	12.0	151	2 A64503	conserved hypothet
14	77.5	11.2	621	2 A75101	aldehyde-ferredoxi
15	77.5	11.2	862	2 T07775	lipoxigenase (EC 1
16	76.5	11.1	6805	2 S20901	titin - rabbit (fr
17	76	11.0	983	2 H64587	cag pathogenicity
18	76	11.0	983	2 F71926	cag pathogenicity
19	75.5	10.9	423	1 VHWVB2	structural polypro
20	75.5	10.9	1245	1 VHWVB2	structural polypro
21	75	10.8	249	2 S75749	hypothetical prote
22	74.5	10.8	625	2 G71072	aldehyde-ferredoxi
23	74.5	10.8	1068	2 F84614	probable kinesin h
24	74	10.7	173	2 S67579	probable membrane
25	73.5	10.6	410	2 C96803	hypothetical prote
26	73	10.5	1098	2 JQ2209	helicase homolog g
27	72.5	10.5	1245	1 VHWVB2	structural polypro
28	72.5	10.5	1878	2 E86189	hypothetical prote
29	72.5	10.5	26926	1 I38344	titin, cardiac mus

30	72	10.4	862	2 S57964	lipoxigenase (EC 1
31	72	10.4	1026	2 T34506	hypothetical prote
32	71	10.3	1324	2 T00386	hypothetical prote
33	70.5	10.2	147	2 S77485	ribosomal protein
34	70	10.1	238	2 AC2485	hypothetical prote
35	69.5	10.0	558	2 F64402	vanadate-sensitive
36	69.5	10.0	585	2 C70341	acetylactate synth
37	69	10.0	343	1 DEBYMP	malate dehydrogena
38	69	10.0	896	2 AB1156	conserved membrane
39	69	10.0	896	2 AE1514	conserved membrane
40	68.5	9.9	289	2 T28311	ORF MSV150 probabl
41	68.5	9.9	357	2 AC3645	flagellar p-ring p
42	68.5	9.9	385	2 T26487	hypothetical prote
43	68.5	9.9	535	2 AF0103	probable sulfatase
44	68.5	9.9	831	2 AB3513	ATPase virB4 homol
45	68.5	9.9	1245	1 VHWVB	structural polypro

ALIGNMENTS

RESULT 1

A60381

allergen Der p II precursor - house-dust mite (Dermatophagoides pteronyssinus)
C;Species: Dermatophagoides pteronyssinus
C;Date: 03-Mar-1993 #sequence_revision 03-Mar-1993 #text_change 09-Jul-2004
C;Accession: A60381

R;Chua, K.Y.; Doyle, C.R.; Simpson, R.J.; Turner, K.J.; Stewart, G.A.; Thomas, W.R.
Int. Arch. Allergy Appl. Immunol. 91, 118-123, 1990

A;Title: Isolation of cDNA coding for the major mite allergen Der p II by IGE plaque in
A;Reference number: A60381; MUID:90256301; PMID:2341191
A;Accession: A60381

A;Status: not compared with conceptual translation

A;Molecule type: mRNA

A;Residues: 1-146 <CHU>

A;Cross-references: UNIPROT:P49278

C;Superfamily: allergen Der p II

F;1-17/Domain: signal sequence #status predicted <SIG>

F;18-146/Product: allergen Der p II #status predicted <MAT>

Query Match 91.8%; Score 635; DB 2; Length 146;

Best Local Similarity 90.7%; Pred. No. 5e-57;

Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVELVPGCHGNPCIIIGRGKPFQLEALFEANONSATAKIEIKASIDG 60

Db 18 DQVDVKDCANHEIKVELVPGCHGSEPCIIHRGKPFQLEAVFEANQNTAKIEIKASIDG 77

QY 61 LSVDPFGIDPNACHYMNCPVNGQQYDIKYTNVPKIAFNSNVVTVKVLGDNGVLACA 120

Db 78 LSVDPFGIDPNACHYMNCPVNGQQYDIKYTNVPKIAFNSNVVTVKVLGDNGVLACA 137

QY 121 IATHAKIRD 129

Db 138 IATHAKIRD 146

RESULT 2

JU0394

allergen Der f II (pFL2) - house-dust mite (Dermatophagoides farinae)

C;Species: Dermatophagoides farinae

C;Date: 30-Sep-1991 #sequence_revision 30-Sep-1991 #text_change 17-Mar-1999

C;Accession: JU0394

R;Yuuki, T.; Okumura, Y.; Ando, T.; Yamakawa, H.; Suko, M.; Haida, M.; Okudaira, H.

Agric. Biol. Chem. 55, 1233-1238, 1991

A;Title: Cloning and expression of cDNA coding for the major house dust mite allergen D

A;Reference number: PS0417; MUID:91291341; PMID:1368682

A;Accession: JU0394

A;Molecule type: mRNA

A;Residues: 1-129 <YUU>

C;Superfamily: allergen Der p II

Query Match 85.5%; Score 592; DB 2; Length 129;

```
Best Local Similarity 82.9%; Pred. No. 9.8e-53;
Matches 107; Conservative 12; Mismatches 10; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVPCGHNPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVMDVDCGSDPCIIHRGKPFQLEALFDANQNTKTAKIEIKASLDG 60

QY 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNVNPVKIAPNSENVVTVKVLGDNGLVACA 120
Db 61 LEIDVPGIDTNACHFMKCPVKGQQYDIKYTNVNPVKIAPKSENENVTVTKLIGDNGVLACA 120

QY 121 IATHAKIRD 129
Db 121 IATHGKIRD 129

RESULT 3
B61241
allergen Der f II precursor - house-dust mite (Dermatophagoides farinae) (fragment)
C:Species: Dermatophagoides farinae
C:Date: 12-May-1994 #sequence_revision 27-Jun-1994 #text_change 13-Sep-1998
C:Accession: B61241; J00395
R:Yuuki, T.; Okumura, Y.; Ando, T.; Yamakawa, H.; Suko, M.; Haida, M.; Dohi, M.; Okudaira
Int. Arch. Allergy Appl. Immunol. 94, 354-356, 1991
A:Title: Synthesis of biologically active recombinant Der f II.
A:Reference number: A61241; MUID:92040281; PMID:1937898
A:Accession: B61241
A:Molecule type: mRNA
A:Residues: 1-138 <YUU>
C:Superfamily: allergen Der p II
F:1-9/Domain: signal sequence (fragment) #status predicted <SIG>
F:10-138/Product: allergen Der f II #status predicted <MAT>

Query Match 85.3%; Score 590; DB 2; Length 138;
Best Local Similarity 82.9%; Pred. No. 1.7e-52;
Matches 107; Conservative 12; Mismatches 10; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVPCGHNPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 10 DQVDVKDCANHEIKVMDVDCGSDPCIIHRGKPFQLEALFDANQNTKTAKIEIKASLDG 69

QY 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNVNPVKIAPNSENVVTVKVLGDNGLVACA 120
Db 70 LEIDVPGIDTNACHFMKCPVKGQQYDAKYTNVNPVKIAPKSENENVTVTKLIGDNGVLACA 129

QY 121 IATHAKIRD 129
Db 130 IATHAKIRD 138

RESULT 4
A61241
allergen Der f II precursor - house-dust mite (Dermatophagoides farinae) (fragment)
C:Species: Dermatophagoides farinae
C:Date: 12-May-1994 #sequence_revision 27-Jun-1994 #text_change 13-Sep-1998
C:Accession: A61241; P50417
R:Yuuki, T.; Okumura, Y.; Ando, T.; Yamakawa, H.; Suko, M.; Haida, M.; Dohi, M.; Okudaira
Int. Arch. Allergy Appl. Immunol. 94, 354-356, 1991
A:Title: Synthesis of biologically active recombinant Der f II.
A:Reference number: A61241; MUID:92040281; PMID:1937898
A:Accession: A61241
A:Molecule type: mRNA
A:Residues: 1-138 <YUU>
A:Note: part of this sequence, including the amino end of the mature protein, was confir
C:Superfamily: allergen Der p II
F:1-9/Domain: signal sequence (fragment) #status predicted <SIG>
F:10-138/Product: allergen Der f II #status experimental <MAT>

Query Match 85.0%; Score 588; DB 2; Length 138;
Best Local Similarity 82.2%; Pred. No. 2.7e-52;
Matches 106; Conservative 13; Mismatches 10; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVPCGHNPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
```

```
Db 10 DQVDVKDCANHEIKVMDVDCGSDPCIIHRGKPFQLEALFDANQNTKTAKIEIKASLDG 69

QY 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNVNPVKIAPNSENVVTVKVLGDNGLVACA 120
Db 70 LEIDVPGIDTNACHFMKCPVKGQQYDIKYTNVNPVKIAPKSENENVTVTKLIGDNGVLACA 129

QY 121 IATHAKIRD 129
Db 130 IATHGKIRD 138

RESULT 5
A61501
allergen Der f II - house-dust mite (Dermatophagoides farinae) (fragment)
C:Species: Dermatophagoides farinae
C:Date: 07-Oct-1994 #sequence_revision 07-Oct-1994 #text_change 09-Jul-2004
C:Accession: A61501
R:Trudinger, M.; Chua, K.Y.; Thomas, W.R.
Clin. Exp. Allergy 21, 33-37, 1991
A:Title: cDNA encoding the major mite allergen Der f II.
A:Reference number: A61501; MUID:91215495; PMID:2021876
A:Accession: A61501
A>Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-129 <TRU>
A:Cross-references: UNIPROT:Q8WQK5
C:Superfamily: allergen Der p II

Query Match 84.5%; Score 585; DB 2; Length 129;
Best Local Similarity 82.2%; Pred. No. 5e-52;
Matches 106; Conservative 12; Mismatches 11; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVPCGHNPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVMDVDCGSDPCIIHRGKPFQLEALFDANQNTKTAKIEIKASLDG 60

QY 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNVNPVKIAPNSENVVTVKVLGDNGLVACA 120
Db 61 LEIDVPGIDTNACHFMKCPVKGQQYDAKYTNVNPVKIAPKSENENVTVTKLIGDNGVLACA 120

QY 121 IATHAKIRD 129
Db 121 IATHAKIRD 129

RESULT 6
S66500
allergen Lep d 1.01 precursor (clone d 1.0102) - Lepidoglyphus destructor
C:Species: Lepidoglyphus destructor
C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 09-Jul-2004
C:Accession: S66500; S48727; S56034
R:Schmidt, M.; Olsson, S.; van der Ploeg, I.; van Hage-Hamsten, M.
FEBS Lett. 370, 11-14, 1995
A:Title: cDNA analysis of the mite allergen Lep d 1 identifies two different isoallergen
A:Reference number: S66499; MUID:95377437; PMID:7649288
A:Accession: S66500
A:Molecule type: mRNA
A:Residues: 1-141 <SCH>
A:Cross-references: UNIPROT:P80384; EMBL:X89014; NID:g999461; PIDN:CAA61419.1; PID:g9994
R:Varela, J.; Ventas, P.; Carreira, J.; Barbaa, J.A.; Gimenez-Gallego, G.; Polo, F.
Eur. J. Biochem. 225, 93-98, 1994
A:Title: Primary structure of Lep d I, the main Lepidoglyphus destructor allergen.
A:Reference number: S48727; MUID:95010146; PMID:7925475
A:Accession: S48727
A:Molecule type: mRNA
A:Residues: 44-141 <VAW>
A:Cross-references: EMBL:X81399; NID:g587449; PIDN:CAA57160.1; PID:g587450
A:Accession: S56034
A:Molecule type: protein
A:Residues: 17-140 <VAR>
A:Note: 53-Asp, 63-Asn, 95-Ile, 104-Asn, 106-Gly and 125-Val were also found
C:Superfamily: allergen Der p II
```


Best Local Similarity 23.7%; Pred. No. 7.8;
Matches 33; Conservative 20; Mismatches 51; Indels 35; Gaps 7;
Qy 1 DOVDVKDCANHEIKEVLVPGCHGNEPCIIGRGKPF-----QLREALFEANQNSA----- 48
Db 211 DKEELKKLSGEAYNDIL-----NAP-----GYFFWKRGQTMAAVEWTNENSALPTNFNS 259
Qy 49 TAKIEIKASIDGLSDVDPGIDPNACHYMNCPNVN-----GQYDIDIKYTNVVPKIAPN-- 100
Db 260 DGSPEFARSIDGYTMEGMKVKQPCYCNMPCGNVVLDAEQSESELDYE-NVALLGANLG 318
Qy 101 ---SENVVVVTKVLGDNGV 116
Db 319 IGKLNVAVLNRIADDMGM 337
RESULT 15
T07775
lipoxigenase (EC 1.13.11.12) LX-3 - potato
C;Species: Solanum tuberosum (potato)
C;Date: 14-May-1999 #sequence_revision 14-May-1999 #text_change 09-Jul-2004
C;Accession: T07775
R;Koloniets, M.V.; Hannapel, D.J.
submitted to the EMBL Data Library, June 1996
A;Reference number: Z16124
A;Accession: T07775
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: mRNA
A;Residues: 1-862 <KOL>
A;Cross-references: UNIPROT:Q43191; EMBL:U60202; NID:gl407704; PIDN:AAB67865.1; PID:gl407704
A;Experimental source: cv. Berolina
C;Genetics:
A;Gene: LX-3
C;Function:
A;Description: catalyzes the oxidation of unsaturated fatty acids with a 1,4-cis,cis per
C;Superfamily: lipoxigenase
C;Keywords: fatty acid oxidation; oxidoreductase
Query Match 11.2%; Score 77.5; DB 2; Length 862;
Best Local Similarity 30.1%; Pred. No. 11;
Matches 31; Conservative 18; Mismatches 35; Indels 19; Gaps 6;
Qy 23 GNEPCIIGRGKPF-----QLREALFEANQNSATAKIEIKASIDGLSDVDPGIDPNACHYMC 78
Db 376 GVNPFVIISRLQEPFPPKSQLDSEVVGQNSITTKHEHIENTLDGLTID-DAIKTNRLYLIN- 433
Qy 79 PLVNGQQYDIKYTNVVPKIAPNSENVVV-----TVKVLGDNGVL 117
Db 434 -----HHDILMPY-VRRRI--NTNTNKLASRTLLFLQDDGTM 467

Search completed: September 9, 2005, 15:23:32
Job time : 26 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 9, 2005, 15:09:49 ; Search time 80 Seconds
(without alignments)
825.728 Million cell updates/sec

Title: US-10-001-245C-36

Perfect score: 692

Sequence: 1 DQVDVDCANHEIKVELVPG.....VLGDNGVLCAIATHAKIRD 129

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Uniprot_03.*

1: uniprot_sprot.*

2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	635	91.8	146	1 ALL2_DERPT	P49278 dermatophag
2	592	85.5	146	1 ALL2_DERFA	Q00855 dermatophag
3	590	85.3	129	2 OBWOF5	Q8WQK5 dermatophag
4	575	83.1	145	1 ALL2_EURMA	Q8TZ22 euroglyphus
5	567.5	82.0	170	2 Q9BIX2	Q9BIX2 dermatophag
6	272.5	39.4	143	1 ALL2_PSOOV	Q96562 psoroptes o
7	251	36.3	125	1 ALL2_GLYDO	Q9NFD4 glycyphagus
8	246.5	35.6	141	1 ALL2_TYRPU	O02380 tyrophagus
9	243	35.1	128	1 AL21_GLYDO	Q9U5P7 glycyphagus
10	240.5	34.8	141	1 ALL2_LEPDS	P80384 lepidoglyph
11	122.5	17.7	163	2 Q7QCK5	Q7QCK5 anopheles g
12	117	16.9	151	2 Q66K95	Q66K95 xenopus tro
13	115.5	16.7	149	1 NPC2_PIG	Q97763 sus scrofa
14	112	16.2	151	1 NPC2_HUMAN	P61916 homo sapien
15	112	16.2	151	1 NPC2_MACFA	P61918 macaca fasc
16	112	16.2	151	1 NPC2_PANTR	P61917 pan troglod
17	106	15.3	150	2 Q6PAR7	Q6PAR7 xenopus lae
18	106	15.3	151	2 Q6NTT7	Q6NTT7 xenopus lae
19	104.5	15.1	149	1 NPC2_CANFA	Q28895 canis famil
20	104	15.0	148	2 Q64FT1	Q64FT1 gekko japon
21	100.5	14.5	149	1 NPC2_BOVIN	P79345 bos taurus
22	95	13.9	158	2 Q7QCK4	Q7QCK4 anopheles g
23	95.5	13.8	154	1 Y146_CABEL	O17271 caenorhabdi
24	95	13.7	149	1 NPC2_MOUSE	Q9Z0J0 mus musculu
25	92.5	13.4	116	2 Q86GB5	Q86GB5 ixodes ric
26	92.5	13.4	155	2 Q7YZR7	Q7YZR7 ixodes ric
27	91.5	13.2	149	2 Q8CHN5	Q8CHN5 rattus norv
28	90.5	13.1	165	2 Q9VH31	Q9VH31 drosophila
29	89.5	12.9	149	1 NPC2_BRARE	Q9DGI3 brachydanio
30	87	12.6	148	1 NPC2_DROME	Q9VQ62 drosophila
31	84.5	12.2	153	2 Q7QCK6	Q7QCK6 anopheles g

32	84.5	12.2	408	2	Q9KBH6	Q9kbh6 bacillus ha
33	83	12.0	151	1	YG27_METJA	Q59021 methanococc
34	82	11.8	159	2	Q9VEN7	Q9ven7 drosophila
35	81	11.7	153	2	Q7QCK8	Q7qck8 anopheles g
36	79.5	11.5	137	2	Q7QQA4	Q7qqa4 anopheles g
37	79.5	11.5	414	2	Q9JHX6	Q9jhx6 mus musculu
38	79	11.4	164	2	Q7PZQ2	Q7pzk2 anopheles g
39	79	11.4	188	2	Q7PZQ3	Q7pzk3 anopheles g
40	79	11.4	422	2	Q8K053	Q8k053 mus musculu
41	79	11.4	711	2	Q8CBC4	Q8cbc4 mus musculu
42	77.5	11.2	214	2	Q6TR70	Q6tr70 pythium aff
43	77.5	11.2	214	2	Q6TR71	Q6tr71 pythium mid
44	77.5	11.2	214	2	Q6TR72	Q6tr72 pythium mon
45	77.5	11.2	273	2	Q72ER8	Q72er8 desulfovibr

ALIGNMENTS

RESULT 1

ALL2_DERPT
ID ALL2_DERPT STANDARD; PRT; 146 AA.
AC P49278;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Mite group 2 allergen Der p 2 precursor (Der p II) (DPX).
GN Names=DERP2;
OS Dermatophagoides pteronyssinus (House-dust mite).
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
OC Acariformes; Sarcoptiformes; Astigmata; Psoroptidia; Analgoidea;
OC Pyroglyphidae; Dermatophagoides.
OX NCBI_TaxID=6956;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90256301; PubMed=2341191;
RA Chua K.Y., Doyle C.R., Simpson R.J., Turner K.J., Stewart G.A.,
Thomas W.R.;
RT "Isolation of cDNA coding for the major mite allergen Der p II by Ige
plaque immunoassay.";
RT Int. Arch. Allergy Appl. Immunol. 91:118-123(1990).
RN [2]
RP SEQUENCE FROM N.A., AND VARIANTS.
RX MEDLINE=21290932; PubMed=11398075; DOI=10.1067/mai.2001.114652;
RA Smith W.-A., Hales B.J., Jarnicki A.G., Thomas W.R.;
RT "Allergens of wild house dust mites: environmental Der p 1 and Der p 2
sequence polymorphisms.";
J. Allergy Clin. Immunol. 107:985-992(2001).
RN [3]
RP PARTIAL SEQUENCE OF 18-57.
RX MEDLINE=89278484; PubMed=2732406;
RA Heymann P.W., Chapman M.D., Aalberse R.C., Fox J.W.,
Platts-Mills T.A.;
RT "Antigenic and structural analysis of group II allergens (Der f II and
Der p II) from house dust mites (Dermatophagoides spp).";
J. Allergy Clin. Immunol. 83:1055-1067(1989).
RN [4]
RP STRUCTURE BY NMR.
RX MEDLINE=98409423; PubMed=9737847; DOI=10.1021/bi980578+;
RA Mueller G.A., Benjamin D.C., Rule G.S.;
RT "Tertiary structure of the major house dust mite allergen Der p 2:
sequential and structural homologies.";
Biochemistry 37:12707-12714(1998).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- ALLERGEN: Causes an allergic reaction in human. Common symptoms of
mite allergy are bronchial asthma, allergic rhinitis and
conjunctivitis.
CC -!- SIMILARITY: Belongs to the NPC2 family.

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CC -----

DR EMBL; AF276239; AAF86462.1; -.
DR PIR; A60381; A60381.
DR PDB; 1A9V; NMR; @=18-146.
DR PDB; 1KTJ; X-ray; A/B=18-146.
DR InterPro; IPR003172; E1_DerP2_DerF2.
DR Pfam; PF02221; E1_DerP2_DerF2; 1.
DR SMART; SM00737; ML; 1.
KW 3D-structure; Allergen; Direct protein sequencing; Polymorphism;
KW Signal.
FT SIGNAL 1 17

FT CHAIN 18 146 Mite group 2 allergen Der p 2.
FT DISULFID 25 136

FT DISULFID 38 44
FT DISULFID 90 95
FT VARIANT 39 39
FT VARIANT 40 40
FT VARIANT 44 44
FT VARIANT 47 47
FT VARIANT 49 49
FT VARIANT 56 56
FT VARIANT 57 57
FT VARIANT 61 61
FT VARIANT 64 64
FT VARIANT 75 75
FT VARIANT 78 78
FT VARIANT 81 81
FT VARIANT 95 95
FT VARIANT 98 98
FT VARIANT 108 108
FT VARIANT 111 111
FT VARIANT 114 114
FT VARIANT 115 115
FT VARIANT 116 116
FT VARIANT 118 118
FT VARIANT 127 127
FT VARIANT 128 128
FT VARIANT 131 131
FT VARIANT 133 133
FT VARIANT 144 144
FT STRAND 19 20
FT STRAND 23 24
FT STRAND 30 34
FT TURN 36 37
FT STRAND 40 40
FT TURN 41 41
FT STRAND 44 47
FT TURN 48 49
FT STRAND 51 59
FT STRAND 64 64
FT STRAND 68 75
FT TURN 76 77
FT STRAND 78 80
FT STRAND 88 88
FT HELIX 89 91
FT STRAND 97 97
FT TURN 99 100
FT STRAND 102 110
FT TURN 113 114
FT STRAND 118 118
FT STRAND 121 129
FT TURN 130 131
FT STRAND 132 139
FT STRAND 142 146
SQ SEQUENCE 146 AA; 15999 MW; 591B2FA7FD26D3AF CRC64;

Query Match 91.8%; Score 635; DB 1; Length 146;
Best Local Similarity 90.7%; Pred. No. 9.5e-55;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0;

QY 1 DOVDVKDCANHEIKVLPVPGCHGNEPCIIIGRKPQLREALFEANONSATAKIEIKASIDG 60
DB 18 DOVDVKDCANHEIKVLPVPGCHGSEPCIIHRGKPFQLEAVFEANQNTKTAKIEIKASIDG 77
QY 61 LSVDPVPGIDPNACHYMNCPLVNGQQYDIKYTNVVKPIAPNSNVVTVKVLGDNGVLACA 120
DB 78 LEVDVPGIDPNACHYMNCPLVNGQQYDIKYTNVVKPIAPNSNVVTVKVLGDNGVLACA 137
QY 121 IATHAKIRD 129
DB 138 IATHAKIRD 146
RESULT 2
ALL2_DERFA
ID ALL2_DERFA STANDARD; PRT; 146 AA.
AC Q00855; P39672; Q26359;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Mite group 2 allergen Der f 2 precursor (Der f II).
GN Name=DERF2;
OS Dermatophagoides farinae (House-dust mite).
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
OC Acariformes; Sarcoptiformes; Astigmata; Psoroptidia; Analgoidea;
OC Pyroglyphidae; Dermatophagoides.
RN NCBI_TaxID=6954;
RX MEDLINE=91291341; PubMed=1368682;
RA Yuuki T., Okumura Y., Ando T., Yamakawa H., Suko M., Haida M.,
RA Okudaira H.;
RT "Cloning and expression of cDNA coding for the major house dust mite
RT allergen Der f II in Escherichia coli.";
RL Agric. Biol. Chem. 55:1233-1238(1991).
RN [2]
RP SEQUENCE OF 4-146 FROM N.A.
RX MEDLINE=94256850; PubMed=8198452;
RA Okuhira H.;
RT "Molecular biology of mite antigens.";
RL Arerugi 43:435-440(1994).
RN [3]
RP DISULFIDE BONDS, AND PARTIAL SEQUENCE.
RX MEDLINE=93283958; PubMed=8508052;
RA Nishiyama C., Yuuki T., Takai T., Okumura Y., Okudaira H.;
RT "Determination of three disulfide bonds in a major house dust mite
RT allergen, Der f II.";
RL Int. Arch. Allergy Immunol. 101:159-166(1993).
RN [4]
RP PARTIAL SEQUENCE OF 18-52.
RX MEDLINE=89278484; PubMed=2732406;
RA Heymann P.W., Chapman M.D., Aalberse R.C., Fox J.W.,
RA Platts-Mills T.A.;
RT "Antigenic and structural analysis of group II allergens (Der f II and
RT Der p II) from house dust mites (Dermatophagoides spp).";
RL J. Allergy Clin. Immunol. 83:1055-1067(1989).
RN [5]
RP STRUCTURE BY NMR.
RX MEDLINE=98079068; PubMed=9417088; DOI=10.1074/jbc.273.1.356;
RA Ichikawa S., Hatanaka H., Yuuki T., Iwamoto N., Kojima S.,
RA Nishiyama C., Ogura K., Okumura Y., Inagaki F.;
RT "Solution structure of Der f 2, the major mite allergen for atopic
RT diseases.";
RL J. Biol. Chem. 273:356-360(1998).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- ALLERGEN: Causes an allergic reaction in human. Common symptoms of
CC mite allergy are bronchial asthma, allergic rhinitis and
CC conjunctivitis.
CC -!- MISCELLANEOUS: The sequence shown here is from clone 2. The N-
CC terminal sequence (AA 1-8) from clone 1 and 11 are not yet known.
CC -!- SIMILARITY: Belongs to the NPC2 family.
CC -----
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DR EMBL; D10447; BAA01239.1; -;
 DR EMBL; D10448; BAA01240.1; -;
 DR EMBL; D10449; BAA01241.1; -;
 DR EMBL; S70378; AAB30829.1; -;
 DR PDB; 1AHM; NMR; @=18-146.
 DR PDB; 1AHM; NMR; @=18-146.
 DR InterPro; IPR003172; E1_DerP2_DerF2.
 DR Pfam; PF02221; E1_DerP2_DerF2; 1.
 DR SMART; SM00737; ML; 1.
 KW 3D-structure; Allergen; Direct protein sequencing; Polymorphism;
 KW Signal.
 FT SIGNAL 1 17
 FT CHAIN 18 146
 FT DISULFID 25 136
 FT DISULFID 38 44
 FT DISULFID 90 95
 FT VARIANT 93 93
 FT VARIANT 105 105
 FT VARIANT 128 128
 FT VARIANT 142 142
 FT CONFLICT 5 8
 FT STRAND 44 46
 FT STRAND 54 54
 FT STRAND 58 60
 FT STRAND 74 75
 FT TURN 76 77
 FT TURN 78 78
 FT TURN 98 99
 FT STRAND 101 103
 FT STRAND 107 107
 FT STRAND 122 128
 FT STRAND 133 139
 FT STRAND 142 144
 SQ SEQUENCE 146 AA; 15802 MW; FAL18206CD88534A CRC64;
 Query Match 85.5%; Score 592; DB 1; Length 146;
 Best Local Similarity 82.9%; Pred. No. 1.7e-50;
 Matches 107; Conservative 12; Mismatches 10; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFLEALFEANQNSATAKIEIKASIDG 60
 DB 18 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFLEALFEANQNTKTAKIEIKASLDG 77
 QY 61 LSVDPVPGIDNPACHYMNCPVNGQQYDIKYTNVPKIAPNSENVVTVKVLGDNGLVACA 120
 DB 78 LEIDVPGIDNACHFMKCPVKGQYDQYDIKYTNVPKIAPNSENVVTVKVLGDNGLVACA 137
 QY 121 IATHAKIRD 129
 DB 138 IATHGKIRD 146

RESULT 3
 Q8WQK5
 ID Q8WQK5 PRELIMINARY; PRT; 129 AA.
 AC Q8WQK5;
 DT 01-MAR-2002 (TrEMBLrel. 20, Created)
 DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Major Der f 2 Isoform (Fragment).
 OS Dermatophagoides farinae (House-dust mite).
 OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
 OC Acariformes; Sarcotiformes; Astigmata; Psoroptidia; Analgoidea;
 OC Pyroglyphidae; Dermatophagoides.
 OX NCBI_TaxID=6954;
 RN [1]

RP SEQUENCE FROM N.A.
 RA Jin H.S., Oh S.H., Hong C.-S.;
 RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AY066008; AAL47677.1; -;
 DR FIR; A61501; A61501.
 DR HSSP; Q00855; 1AHK.
 DR Pfam; PF02221; E1_DerP2_DerF2; 1.
 DR SMART; SM00737; ML; 1.
 FT NON_TER 1 1
 SQ SEQUENCE 129 AA; 14035 MW; 832F72E25FE4F43F CRC64;
 Query Match 85.3%; Score 590; DB 2; Length 129;
 Best Local Similarity 82.9%; Pred. No. 2.3e-50;
 Matches 107; Conservative 12; Mismatches 10; Indels 0; Gaps 0;

QY 1 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFLEALFEANQNSATAKIEIKASIDG 60
 DB 1 DQVDVKDCANHEIKVLPVCGHNEPCIIIGRGKPFLEALFEANQNTKTAKIEIKASLDG 60
 QY 61 LSVDPVPGIDNPACHYMNCPVNGQQYDIKYTNVPKIAPNSENVVTVKVLGDNGLVACA 120
 DB 61 LEIDVPGIDNACHFMKCPVKGQYDQYDIKYTNVPKIAPNSENVVTVKVLGDNGLVACA 120
 QY 121 IATHAKIRD 129
 DB 121 IATHAKIRD 129

RESULT 4
 ALL2_EURMA
 ID ALL2_EURMA STANDARD; PRT; 145 AA.
 AC Q9TZZ2; O96430;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Mite group 2 allergen Eur m 2 precursor.
 GN Name=EURM2;
 OS Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
 OC Acariformes; Sarcotiformes; Astigmata; Psoroptidia; Analgoidea;
 OC Pyroglyphidae; Eukaryophus.
 OX NCBI_TaxID=6958;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 99126275; PubMed=9925958; DOI=10.1159/000024026;
 RA Smith W., Mills K., Hazell L., Hart B.J., Thomas W.;
 RT "Molecular analysis of the group 1 and 2 allergens from the house dust
 mite, Eukaryophus maynei."
 RL Int. Arch. Allergy Immunol. 118:15-22(1999).
 CC 1- SUBCELLULAR LOCATION: Secreted (By similarity).
 CC 1- POLYMORPHISM: The sequence shown is that of isoform Eur m 2.0101.
 CC 1- ALLERGEN: Causes an allergic reaction in human. Common symptoms of
 mite allergy are bronchial asthma, allergic rhinitis and
 conjunctivitis.
 CC 1- SIMILARITY: Belongs to the NPC2 family.
 CC -----
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 CC -----
 DR EMBL; AF047613; AAC82349.1; -;
 DR EMBL; AF047614; AAC82350.1; -;
 DR HSSP; P49278; 1A9V.
 DR InterPro; IPR003172; E1_DerP2_DerF2.
 DR Pfam; PF02221; E1_DerP2_DerF2; 1.
 DR SMART; SM00737; ML; 1.
 KW Allergen; Polymorphism; Signal.
 FT SIGNAL 1 16
 FT CHAIN 17 145
 FT Mite group 2 allergen Eur m 2.

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FT DISULFID 24 135 By similarity.
FT DISULFID 37 43 By similarity.
FT DISULFID 89 94 By similarity.
FT VARIANT 21 I -> V (in Eur m 2 0102).
SQ SEQUENCE 145 AA; 15747 MW; 6655B16C8503A565 CRC64;

Query Match 83.1%; Score 575; DB 1; Length 145;
Best Local Similarity 79.8%; Pred. No. 7.9e-49;
Matches 103; Conservative 14; Mismatches 12; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVELVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 17 DQVDIKDCANHEIKVMVPCGKSEPCVIRGTAFQLEAVFDANQNSNAKIEIKATIDG 76

Qy 61 LSDVDPGIDNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGNGVLACA 120
Db 77 VEIDVPGIDNLLCHFMKCPVKGQYDIKYTNVVPRIAPKSENVVTVKLLGNGVLACA 136

Qy 121 IATHAKIRD 129
Db 137 IATHAKIRD 145

RESULT 5
Q9BIX2 PRELIMINARY; PRT; 170 AA.
AC Q9BIX2; 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Allergen Def f II (Fragment).
OS Dermatophagoides farinae (House-dust mite).
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
OC Acariformes; Sarcotiformes; Astigmata; Psoroptidia; Analgoidea;
OC Pyroglyphidae; Dermatophagoides.
OX NCBI_TaxID=6954;
RN [1]
RP SEQUENCE FROM N.A.
RA Heo M., Xu J., Zhong N.;
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF346905; AAK30133.1; -.
DR HSP; Q00855; IAHK.
DR InterPro: IPR003172; E1_DerP2_DerF2.
DE Pfam; PF02221; E1_DerP2_DerF2; 1.
DR SMART; SM00737; ML; 1.
FT NON_TER 1
FT SEQUENCE 170 AA; 18781 MW; 0C2B58734C9D443A CRC64;

Query Match 82.0%; Score 567.5; DB 2; Length 170;
Best Local Similarity 67.7%; Pred. No. 5.2e-48;
Matches 107; Conservative 12; Mismatches 10; Indels 29; Gaps 1;

Qy 1 DQVDVKDC-----ANHEIKVELVPGCHGNEPCIIGR 31
Db 13 DQVDVKDCGKVCVCFHFFSFLNFKHFLFLVYIHANNEIKVMVDCGCHGSDPCIHR 72

Qy 32 GKPFQLEALFEANQNSATAKIEIKASIDGLSVDPGIDNACHYMCPLVNGQQYDIKYT 91
Db 73 GKPFTEALFDANQNTKATIEIKASIDGLSEIDVPGIDTNACHFMKCPVKGQQYDIKYT 132

Qy 92 WNVPKIAPNSENVVTVKVLGNGVLACATATHAKIRD 129
Db 133 WNVPKIAPNSENVVTVKLLGNGVLACATATHAKIRD 170

RESULT 6
ALL2_PSOOV STANDARD; PRT; 143 AA.
AC Q965E2;
DT 05-JUL-2004 (Rel. 44, Created)
DT 05-JUL-2004 (Rel. 44, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Mite group 2 allergen Pso o 2 precursor (Allergen Pso o A).
```

```
GN Name=ALLA;
OS Psoroptes ovis (Sheep scab mite).
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
OC Acariformes; Sarcotiformes; Astigmata; Psoroptidia; Sarcotoides;
OC Psoroptidae; Psoroptes.
OX NCBI_TaxID=83912;
RN [1]
RP SEQUENCE FROM N.A.
RX PubMed=1191040;
RA Temeyer K.B., Soileau L.C., Pruett J.H.;
RT "Cloning and sequence analysis of a cDNA encoding Pso o II, a mite group II allergen of the sheep scab mite (Acari: Psoroptidae).";
RL J. Med. Entomol. 39:384-391(2002).
RN [2]
RP PARTIAL SEQUENCE OF N-TERMINUS.
RX PubMed=10534947;
RA Pruett J.H.;
RT "Identification and purification of a 16-kDa allergen from Psoroptes ovis (Acarina: Psoroptidae).";
RL J. Med. Entomol. 36:544-550(1999).
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- ALLERGEN: Causes an allergic reaction in human. Common symptoms of mite allergy are bronchial asthma, allergic rhinitis and conjunctivitis.
CC -!- SIMILARITY: Belongs to the NPC2 family.
CC
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CC
CC EMBL; AF187083; AAK61827.1; -.
DR HSP; Q00855; IAHK.
DR InterPro: IPR003172; E1_DerP2_DerF2.
DR Pfam; PF02221; E1_DerP2_DerF2; 1.
DR SMART; SM00737; ML; 1.
KW Allergen; Direct protein sequencing; Signal.
FT SIGNAL 1 17
FT CHAIN 18 143 Mite group 2 allergen Pso o 2.
FT DISULFID 25 134 By similarity.
FT DISULFID 38 43 By similarity.
FT DISULFID 89 94 By similarity.
SQ SEQUENCE 143 AA; 15212 MW; AF03533059DA838D CRC64;

Query Match 39.4%; Score 272.5; DB 1; Length 143;
Best Local Similarity 37.5%; Pred. No. 5.7e-19;
Matches 48; Conservative 35; Mismatches 42; Indels 3; Gaps 3;

Qy 2 QVDVKDCANHEIKVELVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDGL 61
Db 19 KVKFQDCGKGEVESLEVEGCSG-DYCVIHKGKLDLAISVTSNQDSANLKLDIVADINGV 77

Qy 62 SVDVPGIDNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGNGVLACAI 121
Db 78 QIEVPGVDHGGCHYKCPKPKQGHFDVKYTSIPAILPTTKAKII-AKIIGDKLGGC-I 135

Qy 122 ATHAKIRD 129
Db 136 VINGEIQD 143

RESULT 7
ALL2_GLYDO STANDARD; PRT; 125 AA.
AC Q9NFQ4;
DT 05-JUL-2004 (Rel. 44, Created)
DT 05-JUL-2004 (Rel. 44, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Mite group 2 allergen Gly d 2.02.
OS Glyciphagus domesticus (House itch mite).
```

```
CC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;  
CC Acariformes; Sarcophormes; Astigmata; Glyciphagidae; Glyciphagidae;  
CC Glyciphagus.  
CC NCBI_TaxID=105145;  
RX SEQUENCE FROM N.A., AND PARTIAL SEQUENCE OF 1-18.  
RX MEDLINE=21135826; PubMed=11240953; DOI=10.1067/mai.2001.112264;  
RA Gavlin G., Johansson E., Lundin A., Smith A.M., Chapman M.D.,  
RA Benjamin D.C., Derewenda U., van Hage-Hamsten M.;  
RT "Cross-reactivity studies of a new group 2 allergen from the dust mite  
RT Glyciphagus domesticus, Gly d 2, and group 2 allergens from  
RT Dermatophagoides pteronyssinus, Lepidoglyphus destructor, and  
RT Tyrophagus putrescentiae with recombinant allergens.";  
RL J. Allergy Clin. Immunol. 107:511-518 (2001).  
CC -1- SUBCELLULAR LOCATION: Secreted.  
CC -1- ALLERGEN: Causes an allergic reaction in human. Common symptoms of  
CC mite allergy are bronchial asthma, allergic rhinitis and  
CC conjunctivitis.  
CC -1- SIMILARITY: Belongs to the NPC2 family.  
CC -----  
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CC or send an email to license@isb-sib.ch).  
CC -----  
DR EMBL; Y12690; CAA73221.1; --  
DR HSP; Q00855; IAHK.  
DR InterPro; IPR003172; E1_DerP2_DerF2.  
DR Pfam; PF02221; E1_DerP2_DerF2; 1.  
DR SMART; SM00737; ML; 1.  
DR Allergen; Direct protein sequencing; Signal.  
FT SIGNAL 1 15  
FT CHAIN 16 141 Mite group 2 allergen Tyr p 2.  
FT DISULFID 23 132 By similarity.  
FT DISULFID 36 41 By similarity.  
FT DISULFID 87 92 By similarity.  
FT CARBOHYD 103 103 N-linked (GlcNAc...) (Potential).  
SQ SEQUENCE 141 AA; 14851 MW; 38F9520010A04C1 CRC64;  
  
Query Match 35.6%; Score 246.5; DB 1; Length 141;  
Best Local Similarity 39.8%; Pred. No. 2.1e-16;  
Matches 51; Conservative 25; Mismatches 45; Indels 7; Gaps 4;  
  
QY 2 QVVDKCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANONSATAKIEKASIDGL 61  
DB 17 QVRFDCGKKEIASVAVDCEG-DLCVIHKSKEPHVIAEFTANQDTCKIEVKVTGQNL 75  
QY 62 SVDVPGIDPNACHYMNCPVNGQYDIKYTNWPKIAPNSNVVTVKVL--GNGVGLAC 119  
DB 76 EVPPIGLETDCGKVLKCPKLGKPKYTMVNSVNVVVPNIKTV---VKLLATGEHGVGLAC 132  
QY 120 -AIATHAK 126  
DB 133 GAVNTDVK 140  
  
RESULT 9  
ID AL21 GLYDO STANDARD; PRT; 128 AA.  
AC QSU5P7;  
DT 05-JUL-2004 (Rel. 44, Created)  
DT 05-JUL-2004 (Rel. 44, Last sequence update)  
DE 05-JUL-2004 (Rel. 44, Last annotation update)  
DE Mite group 2 allergen Gly d 2.01.  
OS Glyciphagus domesticus (House itch mite).  
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;  
OC Acariformes; Sarcophormes; Astigmata; Glyciphagidae; Glyciphagidae;  
OC Glyciphagus.  
OC NCBI_TaxID=105145;  
RN [1]  
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE OF 1-18.  
RX MEDLINE=21135826; PubMed=11240953; DOI=10.1067/mai.2001.112264;  
RA Gavlin G., Johansson E., Lundin A., Smith A.M., Chapman M.D.,  
RA Benjamin D.C., Derewenda U., van Hage-Hamsten M.;  
RT "Cross-reactivity studies of a new group 2 allergen from the dust mite  
RT Glyciphagus domesticus, Gly d 2, and group 2 allergens from  
RT Dermatophagoides pteronyssinus, Lepidoglyphus destructor, and  
RT Tyrophagus putrescentiae with recombinant allergens.";  
RL J. Allergy Clin. Immunol. 107:511-518 (2001).  
CC -1- SUBCELLULAR LOCATION: Secreted.  
CC -1- ALLERGEN: Causes an allergic reaction in human. Common symptoms of  
CC mite allergy are bronchial asthma, allergic rhinitis and  
CC conjunctivitis.  
CC -1- SIMILARITY: Belongs to the NPC2 family.  
CC -----  
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CC or send an email to license@isb-sib.ch).  
CC -----  
DR EMBL; AJ272216; CAB76459.1; --  
DR InterPro; IPR003172; E1_DerP2_DerF2.  
DR Pfam; PF02221; E1_DerP2_DerF2; 1.  
DR SMART; SM00737; ML; 1.  
KW Allergen; Direct protein sequencing.  
SQ SEQUENCE 125 AA; 13366 MW; 63607D3C6BFOAAE0 CRC64;  
  
Query Match 36.3%; Score 251; DB 1; Length 125;  
Best Local Similarity 42.1%; Pred. No. 6.5e-17;  
Matches 48; Conservative 22; Mismatches 42; Indels 2; Gaps 2;  
  
QY 6 KDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANONSATAKIEKASIDGLSDV 65  
DB 6 KDCGKGEVTELDITDCSG-DPCVTHRGKPLTLEAKFAANQDTTKATIKVLKAGTPIQV 64  
QY 66 PGIDPNACHYMNCPVNGQYDIKYTNWPKIAPNSNVVTVKVLGNGVGLAC 119  
DB 65 PGLTDCGKFKVCKPIKGDPIDFKYTTVPALLPKVK-AEVTAEVLGDHGVGLAC 117  
  
RESULT 8  
ID ALL2 TYRPU STANDARD; PRT; 141 AA.  
AC O02380;  
DT 15-JUL-1998 (Rel. 36, Created)  
DT 15-JUL-1998 (Rel. 36, Last sequence update)  
DT 05-JUL-2004 (Rel. 44, Last annotation update)  
DE Mite group 2 allergen Tyr p 2 precursor.  
OS Tyrophagus putrescentiae (Dust mite).  
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;  
OC Acariformes; Sarcophormes; Astigmata; Acaroidea; Acaridae;  
OC Tyrophagus.  
OC NCBI_TaxID=59818;  
RN [1]  
RP SEQUENCE FROM N.A., AND SEQUENCE OF 16-22.  
RX MEDLINE=98151280; PubMed=9492316;  
RA Eriksson T.L.J., Johansson E., Whitley P., Schmidt M., Elsayed S.,  
RA van Hage-Hamsten M.;  
RT "Cloning and characterization of a group II allergen from the dust  
RT mite Tyrophagus putrescentiae.";  
RL Eur. J. Biochem. 251:443-447 (1998).  
CC -1- SUBCELLULAR LOCATION: Secreted.  
CC -1- ALLERGEN: Causes an allergic reaction in human. Common symptoms of  
CC mite allergy are bronchial asthma, allergic rhinitis and  
CC conjunctivitis.
```

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CC EMBL; AJ249864; CAB59976.1; -.
DR InterPro; IPR003172; E1_Derp2_DerF2.
DR Pfam; PF02221; E1_Derp2_DerF2; 1.
DR SMART; SM00737; ML; 1.
KW Allergen; Direct protein sequencing.
SQ SEQUENCE 128 AA; 13790 MW; 431A02FE89A7B03 CRC64;

Query Match 35.1%; Score 243; DB 1; Length 128;
Best Local Similarity 39.7%; Pred. No. 4.1e-16;
Matches 48; Conservative 32; Mismatches 35; Indels 6; Gaps 4;

QY 2 QVDVKDCANHEIKVLPVGGHNEPCIIIGRKPFQLEALFEALNQNSATKIEIKASIDG- 60
DB 2 KMNFTDCGHNHEIKELSVNCTGNY-CVIRHGKPLTLDAKFDANQDTASVGLVLTALIDG 60
QY 61 LSVDPGIDPNACHMNCPLVNGQQQDIKYTNVVPKIAPNSNVVTVK--VLGDNGVLA 118
DB 61 LAIDIPGLETNACKLMKCPIRKGEHOBLY--NIGSIPDATPRIKAKVKAQLIGEHVLA 118

QY 119 C 119

DB 119 C 119

RESULT 10

ALL2 LEPTS STANDARD; PRT; 141 AA.
AC PR0384; Q8MYK7; Q8MYK8;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Mite group 2 allergen Lep d 2 precursor (Lep d 1) (Lep d I).
OS Lepidoglyphus destructor (Fodder mite).
OC Eukaryota; Metazoa; Arthropoda; Chelicerata; Arachnida; Acari;
OC Acariformes; Tarsozoa; Arthropoda; Chelicerata; Glyciphagidae;
OC Lepidoglyphus.
OX NCBI_TaxID=36936;
RN [1]
RP SEQUENCE FROM N.A. (LEP D 2.0101 AND LEP D 2.0201).
RX MEDLINE=95377437; PubMed=7649288; DOI=10.1016/0014-5793(95)98164-E;
RA Schmidt M., van der Ploeg I., Olsson S., van Hage-Hamsten M.;
RT "cDNA analysis of the mite allergen Lep d 1 identifies two different
RT isoallergens and variants";
RL FEBS Lett. 370:11-14(1995).
RN [2]
RP SEQUENCE FROM N.A. (LEP D 2.0103 AND LEP D 2.0203).
RA Kaiser L., Rasool O., Gavellin G., van Hage-Hamsten M., Johansson E.;
RT "Lep d 2 polymorphisms in wild and cultured Lepidoglyphus mites";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE OF 44-141 FROM N.A., AND SEQUENCE OF 17-140.
RX MEDLINE=95010146; PubMed=7925475;
RA Varela J., Ventas P., Carreira J., Barbas J.A., Gimenez-Gallego G.,
RA Polo F.;
RT "Primary structure of Lep d I, the main Lepidoglyphus destructor
RT allergen";
RL Eur. J. Biochem. 225:93-98(1994).
RN [4]
RP PARTIAL SEQUENCE OF 17-45.
RA Muthiah R., Miller M., Kagen S.;
RT "Barn allergy: isolation and characterization of the major allergens
RT of storage mites: L. destructor";
RL J. Allergy Clin. Immunol. 87:326-326(1991).
RN [5]
RP SEQUENCE OF 17-34.
RX MEDLINE=92362323; PubMed=1355192; DOI=10.1016/0140-6736(92)92152-6;
RA van Hage-Hamsten M., Bergman T., Johansson E., Persson B.,
RA Joernvall H., Haerfast B., Johansson S.G.O.;

RT "N-terminal aminoacid sequence of principal allergen of storage mite
RT Lepidoglyphus destructor.";
RL Lancet 340:614-614(1992).
CC -!- SUBUNIT: Monomer.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- POLYMORPHISM: The sequence shown is that of isoform Lep d 2.0101.
CC -!- ALLERGEN: Causes an allergic reaction in human. Common symptoms of
CC mite allergy are bronchial asthma, allergic rhinitis and
CC conjunctivitis.
CC -!- SIMILARITY: Belongs to the NPC2 family.
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EMBL; X83875; CAA58755.1; -.
DR EMBL; X83876; CAA58756.1; -.
DR EMBL; X89014; CAA61419.1; -.
DR EMBL; AJ487972; CAD32313.1; -.
DR EMBL; AJ487973; CAD32314.1; -.
DR EMBL; X81399; CAA57160.1; -.
DR PIR; S66500; S66500.
DR HSP; Q00855; IAHK.
DR InterPro; IPR003172; E1_Derp2_DerF2.
DR Pfam; PF02221; E1_Derp2_DerF2; 1.
DR SMART; SM00737; ML; 1.
KW Allergen; Direct protein sequencing; Polymorphism; Repeat; Signal.
FT SIGNAL 1 16
FT CHAIN 17 141 Mite group 2 allergen Lep d 2.
FT DOMAIN 64 73 3 X 2 AA repeats of K-V.
FT REPEAT 64 65 1.
FT REPEAT 68 69 2.
FT REPEAT 72 73 3.
FT DISULFID 24 133 By similarity.
FT DISULFID 37 42 By similarity.
FT DISULFID 88 93 By similarity.
FT VARIANT 35 35 T -> S (in Lep d 2.0201 and Lep d
FT VARIANT 48 48 E -> Q (in Lep d 2.0201 and Lep d
FT VARIANT 53 53 E -> D (in Lep d 2.0201 and Lep d
FT VARIANT 63 63 A -> N (in Lep d 2.0201 and Lep d
FT VARIANT 71 71 A -> T (in Lep d 2.0103).
FT VARIANT 90 90 F -> V (in Lep d 2.0201 and Lep d
FT VARIANT 91 91 I -> L (in Lep d 2.0201 and Lep d
FT VARIANT 95 95 V -> I (in Lep d 2.0201 and Lep d
FT VARIANT 104 104 I -> N (in Lep d 2.0201 and Lep d
FT VARIANT 106 106 S -> G (in Lep d 2.0201 and Lep d
FT VARIANT 107 107 G -> M (in Lep d 2.0201 and Lep d
FT VARIANT 116 116 V -> I (in Lep d 2.0201 and Lep d
FT VARIANT 118 118 A -> V (in Lep d 2.0203).
FT VARIANT 125 125 I -> V (in Lep d 2.0201 and Lep d
FT VARIANT 136 136 V -> I (in Lep d 2.0201 and Lep d
FT CONFLICT 26 26 H -> K (in Ref. 5).
FT CONFLICT 30 30 T -> K (in Ref. 5).
SQ SEQUENCE 141 AA; 14773 MW; 9AC96F74D6826FA4 CRC64;
Query Match 34.8%; Score 240.5; DB 1; Length 141;


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Best Local Similarity 36.1%; Pred. No. 8.1e-16;
Matches 44; Conservative 31; Mismatches 44; Indels 3; Gaps 3;

QY 6 KDCANHEIKVLVPGCHGNEPCITGRGKPFQLEALFEANQNSATAKIEIKASIDGLSDV 65
DB 22 KDCGHGEVTELDINGCSG-DTCVLRHGEKMTLEAKFPAQDTAKVTKVLAKVAGTTIQV 80
QY 66 PGIDPNACHYMCPLVNGQQYDIKYTWNVKPIAPNSNVVTVKVLGDNGVLCACAIATHA 125
DB 81 PGLTGDGCKFKCPVKRGEALDFYISGTIPAITPKVK-ADVTABELIGHGVMACG-IVHG 138
QY 126 KI 127
DB 139 QV 140

RESULT 11
Q7QCX5 PRELIMINARY; PRT; 163 AA.
AC Q7QCX5;
DT 01-MAR-2004 (TrEMBLrel. 26, Created)
DT 01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE AGCP1115 (Fragment);
GJ Name=agCG51964; ORFNames=ENSAAGG00000014522;
OS Anopheles gambiae str. PEST.
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Nematocera; Culicoidea; Anopheles.
OX NCBI_TaxID=180454;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PEST;
RA Anopheles Genome Sequencing Consortium;
RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
CC -!- CAUTION: The sequence shown here is derived from an
CC EMBL/GenBank/DBJ whole genome shotgun (WGS) entry which is
CC preliminary data.
DR EMBL; AAB01008859; EAA07711.1; -.
DR InterPro; IPR003172; E1_DerP2_DerF2.
DR Pfam; PF02221; E1_DerP2_DerF2; 1.
FT NON_TER 1
SQ SEQUENCE 163 AA; 17196 MW; 4EAC6C34DD21C04F CRC64;

Query Match 17.7%; Score 122.5; DB 2; Length 163;
Best Local Similarity 30.2%; Pred. No. 0.00043;
Matches 39; Conservative 22; Mismatches 57; Indels 11; Gaps 7;

QY 3 VDVKDCANHE--IKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
DB 34 LEIIQCSNNRPTQEVTVPGC-TSLPCQVPNQSDNFVSFRFAPFTPTILTVDVRSLLG 92
QY 61 LSV--DVPGDPNACHYMN--CLVNGQQYDIKYTWNVKPIAP-NSNVVVTVKVLGDNG 115
DB 93 LFLPYEVEHLRNGCINNINTSCPLTAGQ--SVTLTGTPAVEAPLTGVTVTMEFEITGDGG 150
QY 116 -VLACAIAT 123
DB 151 QVAVCFPAAT 159

RESULT 12
Q66K95 PRELIMINARY; PRT; 151 AA.
AC Q66K95;
DT 25-OCT-2004 (TrEMBLrel. 28, Created)
DT 25-OCT-2004 (TrEMBLrel. 28, Last sequence update)
DT 25-OCT-2004 (TrEMBLrel. 28, Last annotation update)
DE Hypothetical protein.
OS Xenopus tropicalis (Western clawed frog) (Silurana tropicalis).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8364;
```

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RN RP SEQUENCE FROM N.A.
RC TISSUE=Embryo;
RX PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
Altschul S.F., Zebberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Heieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Uedin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
RA Krzywinski M.I., Skaleka U., Smailus D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Matra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Embryo;
RX Klein S., Gerhard D.S.;
RA Submitted (AUG-2004) to the EMBL/GenBank/DBJ databases.
RL EMBL; BC080500; AAH0500.1; -.
DR InterPro; IPR003172; E1_DerP2_DerF2.
DR Pfam; PF02221; E1_DerP2_DerF2; 1.
DR SMART; SM00737; ML: 1.
KW Hypothetical protein.
SQ SEQUENCE 151 AA; 16209 MW; 781FE93CEC4D2D80 CRC64;

Query Match 16.9%; Score 117; DB 2; Length 151;
Best Local Similarity 33.3%; Pred. No. 0.0014;
Matches 31; Conservative 16; Mismatches 40; Indels 6; Gaps 4;

QY 6 KDCANHEIKVL--VPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDGLSV 63
DB 26 KDCGSQSGKLVTLVDVSPC-PEEPCPLVRGSTVTNATFVSNVNSKSASAVVHGIIAGIAV 84
QY 64 DVPGDPNACHY-MNCPLVNGQQYDIKYTWNV 95
DB 85 PPFISEPDGCKSGISCPINSQTY--TYVTKL 115

RESULT 13
NPC2_PIG
ID NPC2_PIG STANDARD; PRT; 149 AA.
AC Q97763;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Epididymal secretory protein E1 precursor (Niemann Pick type C2
DE protein homolog) (16 kDa secretory protein).
GN Name=NPC2;
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
OX NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.; AND SEQUENCE OF 20-39.
RC TISSUE=Epididymis;
RX PubMed=10366780; DOI=10.1016/S1388-1981(99)00070-0;
RA Okamura N., Kiuchi S., Tamba M., Kashima T., Hiramoto S., Baba T.,
RA Dacheux F., Dacheux J.-L., Sugita Y., Jin Y.-Z.;
RT "A porcine homolog of the major secretory protein of human epididymis,
RT HE1, specifically binds cholesterol.";
RL Biochim. Biophys. Acta 1438:377-387 (1999).
CC -!- FUNCTION: May be involved in the regulation of the lipid
```

composition of sperm membranes during the maturation in the
epididymis. Binds cholesterol in a 1:1 ratio.
-!- SUBCELLULAR LOCATION: Secreted.
-!- TISSUE SPECIFICITY: Found in the fluid from the distal caput to
cauda epididymis, not detected in the rete testis and the proximal
and middle caput epididymal fluids.
-!- PTM: N-glycosylated. Found in the epididymal fluid as a 19 kDa
glycoprotein that is processed during its passage through the
epididymis into a 16 kDa protein.
-!- SIMILARITY: Belongs to the NPC2 family.

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or send an email to license@isb-sib.ch).

EMBL; U62253; AAD00096.1; --
DR HSSP; P79345; 1NEP.
DR InterPro; IPR003172; E1_DerP2_DerF2.
DR Pfam; PF02221; E1_DerP2_DerF2; 1.
DR SMART; SM00737; ML; 1.
KW Direct protein sequencing; Glycoprotein; Signal.
FT SIGNAL 1 19
FT CHAIN 20 149 Epididymal secretory protein E1.
FT DISULFID 27 140 By similarity.
FT DISULFID 42 47 By similarity.
FT DISULFID 93 99 By similarity.
FT CARBOHYD 58 58 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 149 AA; 16288 MW; 78F0920057CA0102 CRC64;

Query Match 16.7%; Score 115.5; DB 1; Length 149;
Best Local Similarity 29.7%; Pred. No. 0.0019;
Matches 35; Conservative 21; Mismatches 55; Indels 7; Gaps 5;

QY 1 DQDVVKDCAN--HEIKVLPGCHGNEPCIIGRGKPFQLEALPEANQNSATAKIEIKASI 58
Db 20 EPVHRFDGSGGVGVIKEVNNVPC-PTQPCQLHKGQSYVNVVTPSTQSKGSKAVVHGIV 78
QY 59 DGLSDVDPGIDPNACHY-MNCPLVNGQQYDIKVTNVP-KIAPNSENVVTVVKVLGDN 114
Db 79 MGVPIPPFPDPGCKSGINCPIQKQTY--SYLNKLPVKAEPVPSIKLVVENKLGQDN 134

RESULT 14
ID NPC2_HUMAN STANDARD; PRT; 151 AA.
AC P61916; Q15668; Q29413.
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Epididymal secretory protein E1 precursor (Niemann-Pick disease type
DE C2 protein) (hE1).
GN Name=NPC2;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Epithelium;
RX MEDLINE=93119659; PubMed=8418812;
RT Krull N., Ivell R., Osterhoff C., Kirchoff C.;
RT "Region-specific variation of gene expression in the human epididymis
RT as revealed by in situ hybridization with tissue-specific cDNAs";
RL Mol. Reprod. Dev. 34:16-24(1993).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Ovary;
RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
Hopkins R.P., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
Brownstein M.J., Ustin T.B., Toshiyuki S., Carninci P., Prange C.,
Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
Blakesley R.W., Touchman J.W., Shevchenko Y., Bouffard G.G.,
Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalios D.E.,
Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [3]
RP DISEASE
MEDLINE=20574615; PubMed=11125141; DOI=10.1126/science.290.5500.2298;
RX Naureckiene S., Sleat D.E., Lackland H., Fensom A., Vanier M.T.,
Wattiaux R., Jadot M., Lobel P.;
RT "Identification of HE1 as the second gene of Niemann-Pick C disease";
RL Science 290:2298-2301(2000).
RN [4]
RP VARIANT NP-C2 SER-67.
RX MEDLINE=21473745; PubMed=1567215;
RA Millat G., Chikh K., Naureckiene S., Sleat D.E., Fensom A.H.,
Higaki K., Ellender M., Lobel P., Vanier M.T.;
RT "Niemann-Pick disease type C: spectrum of HE1 mutations and
RT genotype/phenotype correlations in the NPC2 group";
RL Am. J. Hum. Genet. 69:1013-1021(2001).
RN [5]
RP VARIANT NP-C2 MET-39.
RX MEDLINE=22334746; PubMed=12447927; DOI=10.1002/ana.10366;
RA Klunemann H.H., Ellender M., Kaminaki W.E., Snow K., Peyser J.M.,
O'Brien J.F., Munoz D., Schmitz G., Klein H.E., Pendlebury W.W.;
RT "Frontal lobe atrophy due to a mutation in the cholesterol binding
RT protein HE1/NPC2";
RL Ann. Neurol. 52:743-749(2002).
CC -!- FUNCTION: May be involved in the regulation of the lipid
CC composition of sperm membranes during the maturation in the
CC epididymis (By similarity).
CC -!- SUBCELLULAR LOCATION: Secreted (Potential).
CC -!- TISSUE SPECIFICITY: Epididymis.
CC -!- DISEASE: Defects in NPC2 are the cause of Niemann-Pick disease
CC type C2 (NP-C2) [MIM:607625], a fatal autosomal recessive
CC hereditary disease characterized by the accumulation of low-
CC density lipoprotein-derived cholesterol in lysosomes.
CC -!- SIMILARITY: Belongs to the NPC2 family.

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or send an email to license@isb-sib.ch).

EMBL; X67698; CAA47928.1; --
DR EMBL; BC002532; AA02532.1; --
DR EMBL; AL8921; CAA01431.1; --
DR PIR; I38365; I38365.
DR HSSP; P79345; 1NEP.
DR Genew; HGNC:14537; NPC2.
DR MIM; 601015; --
DR MIM; 607625; --
DR InterPro; IPR003172; E1_DerP2_DerF2.
DR InterPro; IPR007110; IG-like.
DR Pfam; PF02221; E1_DerP2_DerF2; 1.
DR SMART; SM00737; ML; 1.

KW Glycoprotein; Polymorphism; Signal.
FT SIGNAL 1 19 Potential.
FT CHAIN 20 151 Epididymal secretory protein E1.
FT DISULFID 27 140 By similarity.
FT DISULFID 42 47 By similarity.
FT DISULFID 93 99 By similarity.
FT CARBOHYD 58 58 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 135 135 N-linked (GlcNAc...) (Potential).
FT VARIANT 39 39 V -> M (in NP-C2).
FT VARIANT 67 67 /FTid=VAR_015948.
FT VARIANT 86 86 S -> P (in NP-C2; dbSNP:11694).
FT VARIANT 86 86 /FTid=VAR_015849.
FT VARIANT 86 86 P -> L (in dbSNP:4688).
FT VARIANT 86 86 /FTid=VAR_011899.
SQ SEQUENCE 151 AA; 16570 MW; B141B611805DC910 CRC64;

Query Match 16.2%; Score 112; DB 1; Length 151;
Best Local Similarity 29.1%; Pred. No. 0.0042;
Matches 37; Conservative 26; Mismatches 50; Indels 14; Gaps 7;

Qy 1 DQVDVKDCANHE--IKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASI 58
Db 20 EPVQFKDCGSDGVKIEVNVSPC-PTQPCQLSKGQSYSVNVVTFSTNQSKSKAVVHGIL 78

Qy 59 DGLSVDPGIDPNACHY-MNCPLVNGQQYDIKTYMNPVKIAPNSE-----NVVVTVKVLGD 113
Db 79 MGVPVPFPPIPEPDGCKSGINCPI----QKDKTYSY-LNKLVPKSEYPSIKLVVWVWQLQDD 133

Qy 114 -NGVLAC 119
Db 134 KNSLFC 140

RESULT 15

NP2 MACFA STANDARD; PRT; 151 AA.
AC P61918; Q15668; Q29413;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Epididymal secretory protein E1 precursor (Niemann Pick type C2
protein homolog) (Epididymal secretory protein 14.6) (ESP14.6).
GN Name=NP2;
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
OX NCBI_TaxID=9541;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Epididymis;
RX MEDLINE=95180740; PubMed=7875608; DOI=10.1016/0378-1119(94)00739-F;
RA Perry A.C.F., Jones R., Hall L.;
RT "The monkey ESP14.6 mRNA, a novel transcript expressed at high levels
in the epididymis.";
RL Gene 153:291-292(1995).
CC -1- FUNCTION: May be involved in the regulation of the lipid
composition of sperm membranes during the maturation in the
epididymis (By similarity).
CC -1- SUBCELLULAR LOCATION: Secreted (Potential).
CC -1- TISSUE SPECIFICITY: Epididymis.
CC -1- SIMILARITY: Belongs to the NP2 family.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; X78134; CAA55013.1; -.
DR PIR; I53929; I53929.

DR HSP; P79345; INEP.
DR InterPro; IPR003172; E1_Derp2_Derp2.
DR InterPro; IPR007110; Ig-like_Derp2.
DR Pfam; PF02221; E1_Derp2_Derp2; 1.
DR SMART; SM00737; ML; 1.
KW Glycoprotein; Signal.
FT SIGNAL 1 19 Potential.
FT CHAIN 20 151 Epididymal secretory protein E1.
FT DISULFID 27 140 By similarity.
FT DISULFID 42 47 By similarity.
FT DISULFID 93 99 By similarity.
FT CARBOHYD 58 58 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 135 135 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 151 AA; 16570 MW; B141B611805DC910 CRC64;

Query Match 16.2%; Score 112; DB 1; Length 151;
Best Local Similarity 29.1%; Pred. No. 0.0042;
Matches 37; Conservative 26; Mismatches 50; Indels 14; Gaps 7;

Qy 1 DQVDVKDCANHE--IKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASI 58
Db 20 EPVQFKDCGSDGVKIEVNVSPC-PTQPCQLSKGQSYSVNVVTFSTNQSKSKAVVHGIL 78

Qy 59 DGLSVDPGIDPNACHY-MNCPLVNGQQYDIKTYMNPVKIAPNSE-----NVVVTVKVLGD 113
Db 79 MGVPVPFPPIPEPDGCKSGINCPI----QKDKTYSY-LNKLVPKSEYPSIKLVVWVWQLQDD 133

Qy 114 -NGVLAC 119
Db 134 KNSLFC 140

Search completed: September 9, 2005, 15:23:02

Job time : 82 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 9, 2005, 15:06:59 ; Search time 27 Seconds
(without alignments)
356.657 Million cell updates/sec

Title: US-10-001-245C-36

Perfect score: 692

Sequence: 1 DQVDVKDCANHEIKVLPV.....VLGDNGVLCAIATHAKIRD 129

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 513545 seqs, 74649064 residues

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued_Patents_AA.*

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2: /cgn2_6/ptodata/1/iaa/5B_COMB.pep.*

3: /cgn2_6/ptodata/1/iaa/6A_COMB.pep.*

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5: /cgn2_6/ptodata/1/iaa/PCTUS_COMB.pep.*

6: /cgn2_6/ptodata/1/iaa/backfiles.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	638	92.2	145	4	US-09-949-889-3
2	635	91.8	145	3	US-08-460-040-6
3	635	91.8	146	1	US-07-945-288-4
4	635	91.8	146	1	US-08-462-831-4
5	635	91.8	146	1	US-08-461-809-4
6	635	91.8	146	1	US-08-461-441-4
7	635	91.8	146	2	US-08-482-142-4
8	635	91.8	146	2	US-08-478-572-4
9	635	91.8	146	5	PCT-US93-08518-4
10	635	91.8	146	5	PCT-US93-08518-4
11	627	90.6	129	1	US-08-462-831-12
12	627	90.6	129	1	US-08-461-809-12
13	627	90.6	129	1	US-08-461-441-12
14	627	90.6	129	5	PCT-US93-08518-12
15	622	89.9	129	1	US-07-945-288-12
16	621	89.7	129	4	US-09-949-889-4
17	603	87.1	129	2	US-08-482-142-157
18	603	87.1	129	3	US-08-478-572-157
19	603	87.1	129	3	US-08-484-296-157
20	596	86.1	129	2	US-08-482-142-159
21	596	86.1	129	3	US-08-478-572-159
22	596	86.1	129	3	US-08-484-296-159
23	592	85.5	129	3	US-08-930-264-4
24	590.5	85.3	130	2	US-08-482-142-158
25	590.5	85.3	130	2	US-08-478-572-158
26	590.5	85.3	130	3	US-08-484-296-158
27	590	85.3	129	3	US-08-930-264-6

28	590	85.3	129	3	US-08-930-264-16	Sequence 16, Appl
29	590	85.3	129	3	US-08-930-264-20	Sequence 20, Appl
30	590	85.3	142	1	US-08-288-888-4	Sequence 4, Appl
31	590	85.3	142	2	US-08-910-075-4	Sequence 4, Appl
32	590	85.3	142	2	US-08-905-801A-4	Sequence 4, Appl
33	589	85.1	129	3	US-08-930-264-18	Sequence 18, Appl
34	588	85.0	129	3	US-08-930-264-2	Sequence 2, Appl
35	588	85.0	142	1	US-08-288-888-2	Sequence 2, Appl
36	588	85.0	142	2	US-08-910-075-2	Sequence 2, Appl
37	588	85.0	142	2	US-08-905-801A-2	Sequence 2, Appl
38	587	84.8	129	3	US-08-930-264-24	Sequence 24, Appl
39	586	84.7	129	3	US-08-930-264-8	Sequence 8, Appl
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41	585	84.5	129	1	US-08-462-831-8	Sequence 8, Appl
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43	585	84.5	129	1	US-08-461-441-8	Sequence 8, Appl
44	585	84.5	129	2	US-08-482-142-8	Sequence 8, Appl
45	585	84.5	129	2	US-08-478-572-8	Sequence 8, Appl

ALIGNMENTS

RESULT 1
US-09-949-889-3
; Sequence 3, Application US/09949889
; Patent No. 6800290
; GENERAL INFORMATION:
; APPLICANT: CONSIGLIO NAZIONALE DELLE RICERCHE
; TITLE OF INVENTION: VARIANTS OF ALLERGENIC PROTEINS OF THE GROUP 2 OF
; FILE REFERENCE: DERMATOPHAGOIDES
; CURRENT APPLICATION NUMBER: US/09/949,889
; CURRENT FILING DATE: 2001-09-12
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: Patent in Ver. 2.1
; SEQ ID NO 3
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-09-949-889-3

Query Match	92.2%	Score	638	DB	4	Length	145
Best Local Similarity	91.5%	Pred. No.	2.1e-67				
Matches	118	Conservative	5	Mismatches	6	Indels	0
						Gaps	0
Qy	1	DQVDVKDCANHEIKVLPVCGHGNPCII	GRGKPFQLEALFEANQNSATAKIKASIDG	60			
Db	17	DQVDVKDCANHEIKVLPVCGHGNPCII	HRGKPFQLEAVFEANQNSATAKIKASIDG	76			
Qy	61	LSVDVPGIDPNACHYMNCPVNGQQYDI	KYTNVPKIDNSNNVVTVKVLGDNGVLACA	120			
Db	77	LEVDPGIDPNACHYMNCPVNGQQYDI	KYTNVPKIDNSNNVVTVKVMGDGVLACA	136			
Qy	121	IATHAKIRD	129				
Db	137	IATHAKIRD	145				
RESULT 2							
US-08-460-040-6							
Sequence 6, Application US/08460040							
Patent No. 6071522							
GENERAL INFORMATION:							
APPLICANT: Thomas, Wayne R.							
TITLE OF INVENTION: Cloning of Mite Allergens							
NUMBER OF SEQUENCES: 8							
CORRESPONDENCE ADDRESS:							
ADDRESSES: LAHIVE & COCKFIELD							
STREET: 60 State Street, suite 510							
CITY: Boston							
STATE: Massachusetts							
COUNTRY: USA							


```
; ATTORNEY/AGENT INFORMATION:
; NAME: MANDRAGOURAS, AMY E.
; REGISTRATION NUMBER: 36,207
; REFERENCE/DOCKET NUMBER: IPC-010CC (IMI-024)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 227-7400
; TELEFAX: (617) 227-5941
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-462-831-4

Query Match          91.8%; Score 635; DB 1; Length 146;
Best Local Similarity 90.7%; Pred. No. 4.7e-67;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0

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       Db         18   DQVDVKDCANHEIKVLVPGCHSGSEPCIIHRGPFPQLEAVFEANKTKAKIEIKASIDG 77

Qy      Qy        61   LSVDPVGIDPNACHYMNCPLNGQQDYDKYTWNVPKIAPNSENVVTVKLVGDNGYLACA 120
       Db         78   LEVDYPGIDPNACHYMKCFLVKGGQYDKYKTWNVPKIAPKSENVVTVKVMGDDGVLACA 137

Qy      Qy        121  IATHAKIRD 129
       Db         138  IATHAKIRD 146
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RESULT 5
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; Sequence 4, Application US/08461809
; Patent No. 5770202
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS FROM
; TITLE OF INVENTION: DERMATOPHAGOIDES
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII TEXT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,809
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/945,288
; FILING DATE: 10 SEPTEMBER 1992
; APPLICATION NUMBER: US 580,655
; FILING DATE: 11 SEPTEMBER 1990
; APPLICATION NUMBER: US 458,642
; FILING DATE: 13 FEBRUARY 1990
; ATTORNEY/AGENT INFORMATION:
; NAME: MANDRAGOURAS, AMY E.
; REGISTRATION NUMBER: 36,207
; REFERENCE/DOCKET NUMBER: IPC-010CC (IMI-024)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 227-7400
; TELEFAX: (617) 227-5941
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:

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; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-461-809-4

Query Match          91.8%; Score 635; DB 1; Length 146;
Best Local Similarity 90.7%; Pred. No. 4.7e-67;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0;

Qy      1  DQVDVKDCANHEIKVELVPGCHGNPECCIIIGRGKPFQLEALPEANQNSATAKIEIKASIDG 60
Db      18  DQVDVKDCANHEIKVELVPGCHGSEPCIIHRGKPFQLEAVPEANQNTKTAKIEIKASIDG 77

Qy      61  LSVDPGIDPNACHMNCPLVNGQQVDIKYTNVVPKIAPNSENVVTVKVLGDNGLVACA 120
Db      78  LEVDVPGIDPNACHMNCPLVNGQQVDIKYTNVVPKIAPNSENVVTVKVMGDDGVLA 137

Qy      121 IATHAKIRD 129
Db      138 IATHAKIRD 146

RESULT 6
US-08-461-441-4
; Sequence 4, Application US/08461441
; Patent No. 5773002
; GENERAL INFORMATION:
; APPLICATION:
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS FROM
; TITLE OF INVENTION: DERMATOPHAGOIDES
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII TEXT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/461,441
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/945,288
; FILING DATE: 10 SEPTEMBER 1992
; APPLICATION NUMBER: US 580,655
; FILING DATE: 11 SEPTEMBER 1990
; APPLICATION NUMBER: US 458,642
; FILING DATE: 13 FEBRUARY 1990
; ATTORNEY/AGENT INFORMATION:
; NAME: MANDRAGOURAS, AMY E.
; REGISTRATION NUMBER: 36,207
; REFERENCE/DOCKET NUMBER: IPC-010CC (IMI-024)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 227-7400
; TELEFAX: (617) 227-5941
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-461-441-4

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QY 1 DQVDVKDCANHEIKEVLVPCGCHNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 18 DQVDVKDCANHEIKKVLVPCGCHSEPCIIHRGKPFQLEAVFEANQNTKTAKIEIKASIDG 77
QY 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLVACA 120
Db 78 LSVDPGIDPNACHYMKCPLVKGQQYDIKYTNVVPKIAPNSENVVTVKVGDDGVLACA 137
QY 121 IATHAKIRD 129
Db 138 IATHAKIRD 146

RESULT 7
US-08-482-142-4
; Sequence 4, Application US/08482142
; Patent No. 5820862
; GENERAL INFORMATION:
; APPLICANT: Garman, Richard
; APPLICANT: Greenstein, Julia
; APPLICANT: Kuo, Mei-chang
; APPLICANT: Rogers, Bruce
; APPLICANT: Franzen, Henry
; APPLICANT: Chen, Xian
; APPLICANT: Evans, Sean
; APPLICANT: Shaked, Ze'ev
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS
; TITLE OF INVENTION: FROM DERMATOPHAGOIDES (HOUSE DUST MITE)
; NUMBER OF SEQUENCES: 207
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: IMMULOGIC PHARMACEUTICAL CORPORATION
; STREET: 610 LINCOLN STREET
; CITY: WALTHAM
; STATE: MA
; COUNTRY: USA
; ZIP: 02154
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII TEXT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/482,142
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/445,307
; FILING DATE: 07 June 1995
; ATTORNEY/AGENT INFORMATION:
; NAME: CRAIG, ANNE I.
; REGISTRATION NUMBER: 32,976
; REFERENCE/DOCKET NUMBER: 017.6US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 466-6000
; TELEFAX: (617) 466-6040
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein

US-08-482-142-4

Query Match 91.8%; Score 635; DB 2; Length 146;
Best Local Similarity 90.7%; Pred. No. 4.7e-67;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0;
QY 1 DQVDVKDCANHEIKEVLVPCGCHNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 18 DQVDVKDCANHEIKKVLVPCGCHSEPCIIHRGKPFQLEAVFEANQNTKTAKIEIKASIDG 77
QY 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLVACA 120
Db 138 IATHAKIRD 146

Db 78 LSVDPGIDPNACHYMKCPLVKGQQYDIKYTNVVPKIAPNSENVVTVKVGDDGVLACA 137
QY 121 IATHAKIRD 129
Db 138 IATHAKIRD 146

RESULT 8
US-08-478-572-4
; Sequence 4, Application US/08478572
; Patent No. 5968526
; GENERAL INFORMATION:
; APPLICANT: Garman, Richard
; APPLICANT: Greenstein, Julia
; APPLICANT: Kuo, Mei-chang
; APPLICANT: Rogers, Bruce
; APPLICANT: Franzen, Henry
; APPLICANT: Chen, Xian
; APPLICANT: Evans, Sean
; APPLICANT: Shaked, Ze'ev
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS
; TITLE OF INVENTION: FROM DERMATOPHAGOIDES (HOUSE DUST MITE)
; NUMBER OF SEQUENCES: 207
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: IMMULOGIC PHARMACEUTICAL CORPORATION
; STREET: 610 LINCOLN STREET
; CITY: WALTHAM
; STATE: MA
; COUNTRY: USA
; ZIP: 02154
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII TEXT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/478,572
; FILING DATE: 07-June-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/445,307
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: CRAIG, ANNE I.
; REGISTRATION NUMBER: 32,976
; REFERENCE/DOCKET NUMBER: 017.6US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 466-6000
; TELEFAX: (617) 466-6040
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-478-572-4

Query Match 91.8%; Score 635; DB 2; Length 146;
Best Local Similarity 90.7%; Pred. No. 4.7e-67;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0;
QY 1 DQVDVKDCANHEIKEVLVPCGCHNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 18 DQVDVKDCANHEIKKVLVPCGCHSEPCIIHRGKPFQLEAVFEANQNTKTAKIEIKASIDG 77
QY 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLVACA 120
Db 78 LSVDPGIDPNACHYMKCPLVKGQQYDIKYTNVVPKIAPNSENVVTVKVGDDGVLACA 137
QY 121 IATHAKIRD 129
Db 138 IATHAKIRD 146


```

RESULT 9
US-08-484-296-4
; Sequence 4, Application US/08484296
; Patent No. 6268491
; GENERAL INFORMATION:
; APPLICANT: Garman, Richard
; APPLICANT: Greenstein, Julia
; APPLICANT: Kuo, Mei-chang
; APPLICANT: Rogers, Bruce
; APPLICANT: Franzen, Henry
; APPLICANT: Chen, Xian
; APPLICANT: Evans, Sean
; APPLICANT: Shaked, Ze'ev
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS
; TITLE OF INVENTION: FROM DERMATOPHAGOIDES (HOUSE DUST MITE)
; NUMBER OF SEQUENCES: 207
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: IMMULOGIC PHARMACEUTICAL CORPORATION
; STREET: 610 LINCOLN STREET
; CITY: WALTHAM
; STATE: MA
; COUNTRY: USA
; ZIP: 02154
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII TEXT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/484,296
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/445,307
; FILING DATE: 07 June 1995
; ATTORNEY/AGENT INFORMATION:
; NAME: CRAIG, ANNE I.
; REGISTRATION NUMBER: 32,976
; REFERENCE/DOCKET NUMBER: 017.6US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 466-6000
; TELEFAX: (617) 466-6040
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-484-296--4

Query Match 91.8%; Score 635; DB 3; Length 146;
Best Local Similarity 90.7%; Pred. No. 4.7e-67;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0;

QY 1 DQDVVKDCANHEIKEVLPVPGCHGNEPCIIIRGKGFQLEALPEANQNSATAKIEIKASIDG 60
DB 18 DQDVVKDCANHEIKKVLVPGCHGSEPCIIIRGKGFQLEALPEANQNTKTAKIEIKASIDG 77
QY 61 LSVDPVPGIDNACHMYNCPLVNGQQYDIKTYTNVPKIPANSENVVTVKVLGDNGLACA 120
DB 78 LEVDVPGIDNACHMYNCPLVKGQQYDIKTYTNVPKIPANSENVVTVKVMGDDGVLACA 137
QY 121 IATHAKIRD 129
DB 138 IATHAKIRD 146

RESULT 10
PCT-US93-08518-4
; Sequence 4, Application PC/TUS9308518
; GENERAL INFORMATION:

```

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; APPLICANT:
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS FROM
; TITLE OF INVENTION: DERMATOPHAGOIDES
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII TEXT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/08518
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/945,288
; FILING DATE: 10 SEPTEMBER 1992
; ATTORNEY/AGENT INFORMATION:
; NAME: MANDRAGOURAS, AMY E.
; REGISTRATION NUMBER: 36,207
; REFERENCE/DOCKET NUMBER: IPC-010CC (IMI-024)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 227-7400
; TELEFAX: (617) 227-5941
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; PCT-US93-08518-4

Query Match          91.8%; Score 635; DB 5; Length 146;
Best Local Similarity 90.7%; Pred. No. 4.7e-67;
Matches 117; Conservative 6; Mismatches 6; Indels 0; Gaps 0

Qy   1  DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLREALFEANQNSATAKIEIKASIDG 60
      |||||
Db   18  DQVDVKDCANHEIKVLVPVPGCHGSEPCIIHRGKPFQLAEAVFEANQNTKTAKIEIKASIDG 77
      |||||

Qy   61  LSVDPVGIDNACHMYNCPLVNGQQYDIKYTNWPKIAPNSENVVVTKVLGDNGVLACA 120
      |||||
Db   78  LEVDVPGIDNACHMYMKCPLVKGGQYDIKYTNWPKIAPKSENVVTVKVMGDDGVGLACA 137
      |||||

Qy   121 IATHAKIRD 129
      |||||
Db   138 IATHAKIRD 146
      |||||

RESULT 11
US-08-462-831-12
; Sequence 12, Application US/08462831
; Patent No. 5552142
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: T CELL EPITOPES OF THE MAJOR ALLERGENS FROM
; TITLE OF INVENTION: DERMATOPHAGOIDES
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk

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RESULT 10
PCT-US93-08518-4
; Sequence 4, Application PC/TUS9308518
; GENERAL INFORMATION:

ADDRESSEE: LAHIVE & COCKFIELD
STREET: 60 STATE STREET, SUITE 510
CITY: BOSTON
STATE: MA
COUNTRY: USA
ZIP: 02109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: ASCII TEXT
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/945,288
FILING DATE: 19920910
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 580,655
FILING DATE: 11 SEPTEMBER 1990
APPLICATION NUMBER: 458,642
FILING DATE: 13 FEBRUARY 1990
ATTORNEY/AGENT INFORMATION:
NAME: MANDRAGOURAS, AMY E.
REGISTRATION NUMBER: P36,207
REFERENCE/DOCKET NUMBER: IPC-010CC (IMI-024)
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 227-7400
TELEFAX: (617) 227-5941
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 129 amino acids
TYPE: AMINO ACID
TOPOLOGY: linear
MOLECULE TYPE: protein
FEATURE:
NAME/KEY: misc feature
LOCATION: 47
OTHER INFORMATION: /label=Xaa is Thr or Ser
FEATURE:
NAME/KEY: misc feature
LOCATION: 113
OTHER INFORMATION: /label=Xaa is Asp or Asn
FEATURE:
NAME/KEY: misc feature
LOCATION: 127
OTHER INFORMATION: /label=Xaa is Ile or Leu
US-07-945-288-12

Query Match 89.9%; Score 622; DB 1; Length 129;
Best Local Similarity 89.1%; Pred.No.1.4e-65;
Matches 115; Conservative 5; Mismatches 9; Indels 0; Gaps 0;
QY 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKKVLVPGCHGSEPCIHRGKPFQLEAVFEANQNKKTAKIEIKASIDG 60
QY 61 LSVDPGIDNACHYMCPLVNGQQYDIKYTWNVPKIAPNSENVVTVKVLDNGVLACA 120
Db 61 LEVDVPGIDNACHYMKCPLVKGQQYDIKYTWNVPKIAPKSENVVTVKVMGXDGVLACA 120
QY 121 IATHAKIRD 129
Db 121 IATHAKIRD 129

Search completed: September 9, 2005, 15:10:16
Job time : 28 secs

	Query Match	100.0%;	Score 692;	DB 14;	Length 129;	
	Best Local Similarity	100.0%;	Pred. No. 4.6e-73;			
	Matches 129; Conservative	0;	Mismatches 0;	Indels 0;	Gaps 0;	
<hr/>						
QY	1	DQDVKDCANHKEIKVLVPCGCHNEPCIIGRGKPFQLEALFEANQNQSATAKIEIKASIDG	60			
Db	1	DQDVKDCANHKEIKVLVPCGCHNEPCIIGRGKPFQLEALFEANQNQSATAKIEIKASIDG	60			
<hr/>						
QY	61	LSVDVEGIDPNACHYMCNPLVNGQQVDIKYTNNVPKIAPNSENVVTVKVLGDNGVLACA	120			

Db 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Qy 121 IATHAKIRD 129
Db 121 IATHAKIRD 129

RESULT 2
US-10-001-245-40
; Sequence 40, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 40
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-40

Query Match 98.8%; Score 684; DB 14; Length 129;
Best Local Similarity 98.4%; Pred. No. 4e-72;
Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Qy 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Db 61 LEVDVPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 3
US-10-001-245-46
; Sequence 46, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 46
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus

US-10-001-245-46

Query Match 98.8%; Score 684; DB 14; Length 129;
Best Local Similarity 98.4%; Pred. No. 4e-72;
Matches 127; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Qy 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Db 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 4
US-10-001-245-42
; Sequence 42, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 42
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-42

Query Match 98.7%; Score 683; DB 14; Length 129;
Best Local Similarity 98.4%; Pred. No. 5.3e-72;
Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKEVLVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Qy 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Db 61 LSVDPGIDPNACHYMNCPVNGQQYDIKYTNWPKIAPNSNVVTVKVLGDSGVLA 120
Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 5
US-10-001-245-44
; Sequence 44, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245

```
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 44
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-44

Query Match          98.7%; Score 683; DB 14; Length 129;
Best Local Similarity 98.4%; Pred. No. 5.3e-72;
Matches 127; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGSEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120
Db 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 6
US-10-001-245-38
; Sequence 38, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; PRIOR FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 38
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-38

Query Match          98.6%; Score 682; DB 14; Length 129;
Best Local Similarity 98.4%; Pred. No. 6.9e-72;
Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120
Db 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 7
US-10-001-245-36
; Sequence 36, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; PRIOR FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 36
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-36

Query Match          98.4%; Score 682; DB 14; Length 129;
Best Local Similarity 98.4%; Pred. No. 6.9e-72;
Matches 127; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120
Db 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 7
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US-10-001-245-48
; Sequence 48, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 48
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-48

Query Match          96.1%; Score 665; DB 14; Length 129;
Best Local Similarity 96.9%; Pred. No. 6.9e-70;
Matches 125; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120
Db 61 LSVDPGIDPNACHMNCPLVNGQQYDIKTYTNVPKIPAPNSENVVTVKVLGDNGLVACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIRD 129

RESULT 8
US-10-001-245-52
; Sequence 52, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001,245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 52
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-52

Query Match          94.9%; Score 657; DB 14; Length 129;
Best Local Similarity 95.3%; Pred. No. 6e-69;
Matches 123; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
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; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001.245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 50
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-50

Query Match          94.7%; Score 655; DB 14; Length 129;
Best Local Similarity 95.3%; Pred. No. 1e-68;
Matches 123; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLACA 120
Db 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIQD 129

RESULT 13
US-10-001-245-94
; Sequence 94, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001.245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 94
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-94

Query Match          93.6%; Score 648; DB 14; Length 129;
Best Local Similarity 93.8%; Pred. No. 6.9e-68;
Matches 121; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLACA 120
Db 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIRD 129

Query Match          93.4%; Score 646; DB 14; Length 129;
Best Local Similarity 93.0%; Pred. No. 1.2e-67;
Matches 120; Conservative 3; Mismatches 6; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
Db 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60

Qy 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLACA 120
Db 61 LSVDPGIDPNACHYMCPLVNGQQYDIKYTNVVPKIAPNSENVVTVKVLGDNGLACA 120

Qy 121 IATHAKIRD 129
Db 121 IATHAKIRD 129

RESULT 14
US-10-001-245-171
; Sequence 171, Application US/10001245
; Publication No. US20030175312A1
; GENERAL INFORMATION:
; APPLICANT: HOLM, Jens
; APPLICANT: IPSEN, Henrik
; APPLICANT: LARSEN, Jorgen N.
; APPLICANT: SPANGFORT, Michael D.
; TITLE OF INVENTION: No. US20030175312A1 mutant allergens
; FILE REFERENCE: 4305/1H942-US2
; CURRENT APPLICATION NUMBER: US/10/001.245
; CURRENT FILING DATE: 2001-11-15
; PRIOR APPLICATION NUMBER: US 60/298,170
; PRIOR FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/249,361
; PRIOR FILING DATE: 2000-11-16
; NUMBER OF SEQ ID NOS: 217
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 171
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Dermatophagoides pteronyssinus
US-10-001-245-171

Query Match          92.9%; Score 643; DB 14; Length 129;
Best Local Similarity 92.2%; Pred. No. 2.7e-67;
Matches 119; Conservative 4; Mismatches 6; Indels 0; Gaps 0;

Qy 1 DQVDVKDCANHEIKVLPVPGCHGNEPCIIGRGKPFQLEALFEANQNSATAKIEIKASIDG 60
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Db	1	QVDDVKDCANHEIKKVLVPGCHGSEPCIIHRGKPFQLEAVFEANQNSKTAKIEIKASIDG	60
Qy	61	LSVDVPGIDPNACHYNNCPVLVNGQQYDIKYTNVVPKIAPNSENWVTVKVLGDNGVLACA	120
Db	61	LEVDPGIDPNACHYMKCPLVKGGQYDIKYTNVVPKIAPKSENWVTVKVGDNGLACA	120
Qy	121	IATHAKIRD	129
Db	121	IATHAKIRD	129

Search completed: September 9, 2005, 15:20:09

Job time : 393 secs